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FINAL REPORT

STUDIES OF THE FUNDAMENTALS OF SOLID STATE BATTERIES

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- 4) "Polyphosphazene-Poly(olefin oxide) Mixed Polymer Electrolytes. II. Characterization of MEEP/PPO-(LiX)_n", J. Electrochem. Soc., in press.
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**Dimensionally Stable MEEP-Based Polymer Electrolytes
and Solid-State Lithium Batteries**

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ABSTRACT

Several methods have been developed to dimensionally stabilize polymer electrolytes based on poly[bis-(methoxyethoxyethoxide)phosphazene, known as MEEP. In contrast to the poor dimensional stability exhibited by complexes of MEEP with most Li salts, those prepared with LiAlCl_4 have been isolated as the first example of free-standing $\text{MEEP}-(\text{LiX})_n$ films. The mechanical properties of dimensionally unstable $\text{MEEP}-(\text{LiX})_n$ complexes can be significantly improved by forming composites with polymers such as poly(ethylene oxide), poly(propylene oxide), poly(ethylene glycol diacrylate) and poly(vinyl pyrrolidinone). The conductivity of $6.7 \times 10^{-5} \text{ ohm}^{-1} \cdot \text{cm}^{-1}$ at 25°C exhibited by 55 w/o MEEP/45 w/o PEO- $[\text{LiN}(\text{CF}_3\text{SO}_2)_2]_{0.13}$ is among the highest values reported to date for a dimensionally stable electrolyte. The preparation, and conductivity, calorimetric and electrochemical studies of these electrolytes are described. Cyclic voltammetric data indicated that these electrolytes have anodic stability at least up to 4.5V versus Li^+/Li . They have shown excellent compatibility with Li metal making them suitable for use as Li^+ conductive solid electrolytes in solid-state Li batteries. Li/TiS_2 solid-state cells utilizing some of these electrolytes have exceeded 200 cycles.

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INTRODUCTION

Li^+ -conductive polymer electrolytes derived from Li salt complexes of poly[bis(methoxyethoxyethoxide)phosphazene], or MEEP, due to their high ambient temperature conductivity, are of considerable importance for the fabrication of solid-state Li batteries⁽¹⁻³⁾. They have exhibited 3-4 orders of magnitude higher conductivity at room temperature than electrolytes based on their poly(ethylene oxide), (PEO), counterparts. However, the poor mechanical properties of MEEP-based electrolytes has presented practical problems when attempts were made to fabricate all-solid-state Li batteries incorporating them^(2,4). At room temperature and above, these complexes are glutinous materials with a tendency to flow under pressure. Several approaches have been pursued to enhance the mechanical properties of MEEP. These include chemical crosslinking of MEEP with poly(ethylene glycol)⁽⁵⁾, irradiation of MEEP or $\text{MEEP}-(\text{LiX})_{0.25}$ complexes with various doses of ^{60}Co γ -ray to induce crosslinking^(6,7), and the use of porous, fiberglass support-matrix⁽⁴⁾.

We have found^(2,4,8,9) that dimensional stability of $\text{MEEP}-(\text{LiX})_n$ complexes can be improved to prepare free-standing films by forming composites with high molecular weight PEO or poly(propylene oxide), PPO. Films were cast from solutions containing requisite amounts of MEEP, PEO or PPO, and a Li salt in acetonitrile. We have now extended the class of dimensionally stable MEEP-based composite polymer electrolytes to include blends of MEEP with *in situ* photopolymerized poly(ethylene glycol diacrylate) (PEGDA) and poly(vinyl pyrrolidinone) (PVP). Although the latter two polymers are not poly(olefin oxides), composite electrolytes formed between $\text{MEEP}-(\text{LiX})_n$ and each of them have shown electrical and mechanical characteristics similar to those containing the poly(olefin oxide)s.

We have also discovered that complexes of MEEP and LiAlCl_4 can be isolated as dimensionally stable electrolytes. To our knowledge, this is the first example of a MEEP-(LiX)_n electrolyte which can be processed into free-standing, thin films without such physical treatment as γ -radiation or the addition of a mechanically strong second polymer. Not only are the LiAlCl_4 complexes mechanically stable, but they also possess conductivity very close to that of other MEEP-(LiX)_n electrolytes.

The electrical, thermal and structural properties of these various dimensionally stable electrolytes we have prepared are reported here. While the emphasis in this paper is on $\text{MEEP-(LiAlCl}_4)_n$ and the composite electrolytes formed with PVP and PEGDA, some of the results for MEEP/PEO-(LiX)_n and MEEP/PPO-(LiX)_n composites are included to put the structural properties and electrical behavior of the general family of dimensionally stable MEEP-based electrolytes into perspective. In addition, solid-state Li/TiS_2 cells using a selected group of these electrolytes have been constructed and electrochemically cycled at different temperatures and rates.

EXPERIMENTAL

MEEP was synthesized according to the procedure of Allcock et al.⁽¹⁰⁾ The details of our preparation have been reported earlier⁽⁹⁾.

PEO (Polysciences) of average molecular weight 5×10^6 was used as-received after drying in vacuum at 50°C . LiClO_4 , LiBF_4 and $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ were dried in vacuum at temperatures around 120°C . LiAlCl_4 obtained from Anderson Physics was used as received. Tetraethylene glycol diacrylate (Aldrich), the precursor for PEGDA films, was purified by passing it through an inhibitor remover column. Vinyl pyrrolidinone (Aldrich) was purified by distilling under vacuum.

Pure MEEP-(LiAlCl₄)_n electrolytes were prepared by dissolving MEEP and the corresponding amount of LiAlCl₄ in acetonitrile. The mixture was then homogenized by stirring in order to prepare bubble-free electrolyte films. The films, cast by pouring the solution onto Teflon dishes, were allowed to dry at room temperature inside an argon-filled glove box. Subsequently, they were dried under vacuum (<0.1 Torr) at 45-50°C for 2 days. The dried films, thickness of which varied between 4 and 6 mils, were stored inside the glove-box. The preparation of the composite electrolytes was carried out in two different ways. For the MEEP/PEO-(LiX)_n and MEEP/PPO-(LiX)_n electrolytes, appropriate amounts of PEO or PPO, MEEP and a Li salt were dissolved together in acetonitrile and cast into thin films as described above. Dimensional stabilization of MEEP-electrolytes by reinforcement with PVP or PEGDA was carried out in the following manner. A known quantity of vinyl pyrrolidinone or tetraethylene glycol diacrylate in conjunction with a few drops of a photoinitiator was added to a solution of MEEP-(LiClO₄)_{0.25} in acetonitrile. After evaporating off acetonitrile at room temperature, the mixture was irradiated with UV-light (λ=365 nm) for about five minutes, which resulted in the formation of mechanically stable films. They were again dried at 45-50°C under vacuum for 2 days.

The electrochemical behavior of neat MEEP-(LiClO₄)_{0.25} electrolytes was studied after stabilizing them in a separator-matrix (4). It was accomplished by spreading the electrolyte on a 2 mil thick fiberglass separator paper and wetting it with a few drops of tetrahydrofuran until the electrolyte was homogeneously distributed within the porous matrix. The solvent was then pumped off.

Characterization of the polymer electrolytes was carried out using complex impedance spectroscopy, differential scanning calorimetry (DSC), cyclic voltammetry, X-ray diffraction and infrared spectroscopy. The DSC data were obtained using a

Perkin-Elmer DSC7-calorimeter. Complex impedance measurements between 100 kHz and 5 Hz were performed using an EG&G Par Model M378 impedance system and stainless steel electrodes⁽⁸⁾. X-ray diffraction patterns were obtained using a Rigaku X-ray diffractometer and $\text{CuK}\alpha$ radiation. Infrared spectra were recorded on an IBM 9000 FTIR spectrometer with the electrolyte sandwiched between KBr windows.

Cyclic voltammetry on the polymer electrolytes was carried out at 75°C on a stainless steel or a gold working electrode using Li both as the counter and reference electrodes. An EG&G PAR Model 175 Universal Programmer in conjunction with an ECO Model 551 potentiostat and a Bascom-Turner X-Y Recorder (Model 4110) was used to carry out the voltammetric experiments. The electrolyte was sandwiched between the working and the Li electrodes.

Transport number for Li^+ in the polymer electrolytes was determined using the potentiostatic polarization technique in cells of the configuration Li/polymer electrolyte/Li⁽¹¹⁾. A constant voltage of 25 mV was applied to the cell. The resulting current was monitored as a function of time with a recorder until the steady state current was attained. The transport number, t_+ , was calculated as the ratio between the values of the steady-state and initial currents.

Experimental Li/TiS_2 cells were constructed with the polymer electrolyte sandwiched between a 0.005 cm thick Li foil anode pressed onto a Ni screen, and a TiS_2 composite cathode. The TiS_2 electrodes were prepared by spreading a mixture of 80 weight-percent (w/o) TiS_2 , having a stoichiometric composition of $\text{T}_{1.02}\text{S}_2$ prepared in-house, and 20 w/o polymer electrolyte of the composition 70 MEEP/30 $\text{PEO-(LiX)}_{0.13}$ on one side of a Ni foil. All the cells were constructed in the cathode limited configuration. The electrode package

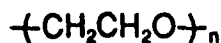
was secured between plastic plates with a positive pressure applied to the plates by means of compression springs to allow good interfacial contact between the electrodes. Cells were cycled between 3.0 and 1.7V at constant currents and at several temperatures.

RESULTS AND DISCUSSION

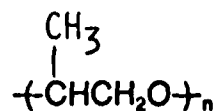
Several dimensionally stable, Li^+ conductive MEEP-based polymer electrolytes have now been prepared in our laboratories, allowing the utilization of these highly conductive electrolytes in all-solid-state rechargeable Li batteries which can be operated at ambient temperatures. The family of dimensionally stable electrolytes based on MEEP now includes:

- MEEP- $(\text{LiAlCl}_4)_n$ complexes.
- MEEP/PEGDA- $(\text{LiX})_n$ or MEEP/PVP- $(\text{LiX})_n$ electrolytes, where LiX include LiBF_4 and LiClO_4 obtained by *in situ* photopolymerization.
- MEEP/PEO- $(\text{LiX})_n$ or MEEP/PPO- $(\text{LiX})_n$ complexes, where LiX include LiBF_4 , LiClO_4 , LiAsF_6 , LiCF_3SO_3 , LiAlCl_4 and $\text{LiN}(\text{SO}_2\text{CF}_3)_2$.
- MEEP- $(\text{LiX})_n$ supported on a porous matrix such as a fiberglass separator.

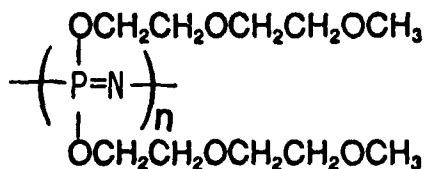
The structural formulas of the polymers employed in this study are given below and are useful for visualizing the structure-property relationships of the related electrolytes.



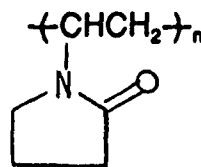
PEO



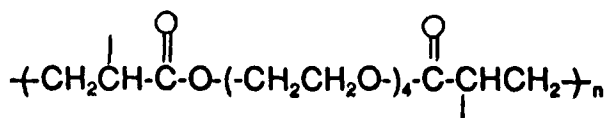
PPO



MEEP



PVP



PEGDA

While PVP is shown as a linear polymer, it is possible that a certain amount of crosslinking does occur in the photopolymerization process. The PEGDA is believed to be a predominantly crosslinked polymer⁽¹²⁾.

Explanations for formulas of the electrolytes described in this paper are as follows. In the single polymer electrolytes, PEO-(LiX)_n, PPO-(LiX)_n and MEEP-(LiX)_n, the subscript n is the Li⁺ to O mole ratio wherein complexes of the stoichiometries (PEO)₆LiX, (PPO)₆LiX and (MEEP)₄LiX are assumed to form. In the mixed polymer electrolytes the polymer ratios are given in weight-percentages. The concentration of the Li salt in the MEEP/PEO and MEEP/PPO mixed electrolytes is selected in such a way that the Li⁺ to O mole ratios are additive functions of those in the corresponding single polymer electrolytes. In the MEEP/PVP and MEEP/PEGDA electrolytes only MEEP is assumed to complex with Li⁺.

Physical Properties of MEEP-Based Electrolytes: MEEP and its Li salt complexes such as MEEP-(LiBF₄)_{0.25}, MEEP-(LiClO₄)_{0.25} and MEEP-(LiSO₃CF₃)_{0.25} are glutinous materials with a tendency to flow even under slight compression. Against this experience, it was surprising to find that MEEP doped with LiAlCl₄ could be processed into dimensionally stable, free-standing thin films. As mentioned earlier, this is the first example of a MEEP-(LiX)_n complex which demonstrates such property without the addition of a second polymer or immobilization in a support matrix or the use of high energy radiation to crosslink. Homogeneous and uniform films of complexes having various MEEP to LiAlCl₄ ratios have been prepared and studied. Thin films of these electrolytes appeared almost clear. These films were found to be highly sensitive to moist atmosphere. For example, a piece of film lost all its dimensional stability and became a liquid mass upon sitting in the normal atmosphere overnight.

MEEP-based electrolytes containing a reinforcement of PEGDA or PVP also exhibited excellent dimensional stability and film homogeneity. These films while appeared as transparent as MEEP-(LiAlCl₄)_n, were softer than the latter. Prolonged UV irradiation for a period of 20 minutes or more, instead of the typical 5 minutes, led these polymer composite films to acquire a yellow coloration.

Mechanically stable electrolyte films could also be cast from solutions of a mixture of MEEP and PEO or MEEP and PPO in acetonitrile containing an appropriate Li salt. It appears that when the electrolyte matrix is interspersed with an appropriate amount of PEO or PPO, a dimensionally stable film can be prepared irrespective of the salt. We have demonstrated this with LiBF₄, LiClO₄, LiAsF₆, LiCF₃SO₃, LiAlCl₄ and LiN(SO₂CF₃)₂. An important observation was that when the mixture contained MEEP and PEO, a homogeneous electrolyte could be prepared only if the amount of MEEP was ≤ 70 weight percent

of the polymer mixture. For higher MEEP/PEO weight ratios, the films appeared heterogeneous with isolated regions of PEO and MEEP. Mixtures of MEEP and PPO, on the other hand, formed homogeneous solid solutions at all proportions, and mechanically strong MEEP/PPO-(LiX)_n electrolytes could be prepared if the amount of PPO was at least 20 w/o of the polymer mixture. Some of our results on MEEP/PEO-(LiX)_n and MEEP/PPO-(LiX)_n have already been reported^(8,9).

Structure of MEEP-Based Electrolytes: We may begin our discussion of the structure of these electrolytes by recalling that polymer electrolytes are complexes of Li salts with polymers having electron donor atoms such as O or N. Lithium salts will dissolve in the polymer if the free-energy for complex formation between Li⁺ and the polymer exceeds the lattice energy of the salt. The ionic conductivity of the resulting electrolyte will depend on a number of factors including the permittivity of the medium, and the concentration of the ions and their mobility. The ionic conductivity of polymer electrolytes is believed to be through the disordered (amorphous) domains of the polymer matrix and is assisted by the large-amplitude segmental motion of the polymer backbone. The structures of PEO-(LiX)_n complexes and their ionic conductivity mechanisms have been discussed^(13,14).

X-ray analysis revealed that all of the MEEP-based electrolytes, except the MEEP/PEO-(LiX)_n composites where LiX = LiClO₄ and LiBF₄, lacked Bragg peaks characteristic of crystalline materials. These observations are also consistent with the differential scanning calorimetric (DSC) data discussed below. X-ray patterns for several MEEP/PEO-(LiClO₄)_n complexes are presented in Table 1 and are compared to the patterns for LiClO₄, PEO and PEO-(LiClO₄)_{0.13}. The data for the latter electrolyte show peaks due to PEO at 2.32Å and 1.93Å along with a strong line at 3.00Å, most probably due to a crystalline PEO-(LiClO₄)_n. Crystalline PEO-(LiClO₄)_n complexes have been previously

identified⁽¹³⁾, including a material in which $n = \sim 0.33$ melting at about 160°C, and a complex in which $n = \sim 0.17$ with a melting point of about 65°C. On the basis of the Li salt to polymer ratio the complex in our case most probably is the latter. The X-ray diffraction patterns of MEEP/PEO-(LiClO₄)_{0.13} complexes show the principal line of PEO at 1.96Å and PEO-(LiClO₄)_n at 3.02Å. The line at 2.02Å is common to all of the MEEP/PEO mixed electrolytes and its intensity increases with the amount of MEEP in the sample. Since MEEP-(LiClO₄)_n itself is amorphous, we may ascribe the line at 2.02Å to a mixed complex formed between LiClO₄ and, MEEP and PEO. X-ray data for MEEP/PEO-(LiBF₄)_n could also be interpreted in a similar way. Thus, the MEEP/PEO-(LiX)_{0.13} mixed electrolyte films appear to be a composite of the three crystalline phases of PEO, PEO-(LiX)_n, MEEP/PEO-(LiX)_n plus amorphous MEEP and possibly an amorphous MEEP-(LiX)_n complex. By analogy, a similar distribution of single and mixed polymer complexes can be visualized in the case of MEEP/PPO-(LiX)_n composites. However, X-ray data indicated the latter to be amorphous⁽⁹⁾. It may be argued that the MEEP/PVP-(LiX)_n and MEEP/PEGDA-(LiX)_n are also composed of single and mixed polymer complexes of Li salts, although X-ray and DSC data were of little help in resolving this issue. Both of these complexes were amorphous. The MEEP/PEO mixed complexes of LiN(SO₂CF₃)₂, in contrast to the electrolytes derived from the other salts, were fully amorphous. As shown later in this paper, they are among the highest conducting dimensionally stable electrolytes reported to date. It should also be noted that MEEP-(LiAlCl₄)_n despite its dimensional stability was amorphous to X-rays in the same manner as the glutinous complexes.

Infrared spectra were examined to gain further insight into the structures of the electrolytes. The IR spectra for PEO, PEO-(LiClO₄)_{0.13}, MEEP, MEEP-(LiClO₄)_{0.25}, and

MEEP-(LiAlCl₄)_{0.25} are presented in Figures 1 and 2. The 1500-500 cm⁻¹ spectral region was the most informative. The infrared spectra of PEO and a number of PEO-(LiX)_n complexes have been discussed by Papke et al.⁽¹⁵⁾, although their paper did not include LiClO₄ complexes. Also, Papke et al.'s PEO had a molecular weight of 6x10⁵ as opposed to 5x10⁶ for the material in this work, and they primarily dealt with PEO-(LiX)_{0.25} while the focus here is on PEO-(LiX)_{0.125}. The infrared spectrum of our PEO is identical to that reported in ref. 15. However, in several respects our spectrum of PEO-(LiClO₄)_{0.125} differs from that of PEO-(LiBF₄)_{0.25} and other complexes given in ref. 15. Relative to the spectrum of PEO, our spectrum of PEO-(LiClO₄)_{0.125} in Figure 1 reveals clearly discernible new peaks at 1590, 1371, 1320, 920, 827, 815, 625, and 572 (doublet) cm⁻¹. The broad, apparently triplet, peak around 1110 cm⁻¹ in PEO has become broader in PEO-(LiClO₄)_{0.125}. The 625 cm⁻¹ peak in the complex is due to ClO stretching vibrations of the ClO₄⁻ anion and the broadening of the C-O-C linkage vibration around 1110 cm⁻¹ is indicative of C-O → Li⁺ complexation. The presence of the new IR peaks in addition to the peaks for neat PEO in our complex is consistent with a material containing both uncomplexed PEO and the PEO-(LiClO₄)_n complex. A very distinct feature of the spectrum is the doublet at 572 cm⁻¹ associated with O-C-C bending in the PEO-(LiClO₄)_{0.125} complex, a counterpart of the peak at 520 cm⁻¹ in neat PEO. (This vibrational assignment is according to ref. 15). It appears that Papke et al. did not observe a clear distinction between PEO and PEO-(LiX)_n because in their complexes n was about 0.25 so that very little free PEO existed.

The IR spectra of MEEP, MEEP-(LiAlCl₄)_{0.25} and MEEP-(LiClO₄)_{0.25}, depicted in Figure 2, are characterized by a number of broad absorption peaks in the 750-1500 cm⁻¹ region from which no significant information relevant to the structure of the complex could be gained. In these materials the P=N and C-O-C stretching vibrations are found at about

1250 and 1100 cm^{-1} , respectively. The Cl-O stretching vibrational peak in the LiClO_4 complex is found at 623 cm^{-1} . This peak was also present in the $\text{MEEP/PEO-(LiClO}_4\text{)}_{0.13}$ complexes. The IR spectrum of $\text{MEEP-(LiAlCl}_4\text{)}_{0.25}$ shows features similar to those found in $\text{MEEP-(LiClO}_4\text{)}_{0.25}$. A peak due to AlCl_4^- is not observed since its absorption is found around 480 cm^{-1} .⁽¹⁶⁾

Differential scanning calorimetric (DSC) traces of MEEP-(LiX)_n , where $\text{LiX} = \text{LiClO}_4$, LiBF_4 , LiSO_3CF_3 , were all without any peaks in the 25 to 200°C range. This confirms their amorphous characteristics. The DSC traces of three $\text{MEEP-(LiAlCl}_4\text{)}_n$ films are presented in Figure 3. While the general profile is again in agreement with the electrolyte being amorphous, a broad endotherm extending from around 60°C to 140°C is observed. The origin of the endotherm is presently not known; it may correspond to the removal of the solvent which remains in the film after being cast from solution. Alternately, the electrolyte may be partially crystalline with the crystallinity distributed over microdomains so as to be unresponsive to X-rays. The broad endotherm may correspond to the ill-defined melting of the crystalline phase.

The DSC traces of all the composites, except MEEP/PEO-(LiX)_n , confirmed them to be amorphous materials. The DSC data for 55 $\text{MEEP/45 PEO-(LiClO}_4\text{)}_{0.13}$ and $\text{PEO-(LiClO}_4\text{)}_{0.13}$ are presented in Figure 4. The latter shows two closely lying peaks at 58 and 65°C, probably associated with the melting of crystalline PEO and a $\text{PEO-(LiClO}_4\text{)}_n$ complex. The DSC thermogram for 55 $\text{MEEP/45 PEO-(LiX)}_n$ appears to contain melting point transitions at 53 and 63°C with possibly a third, lower melting, peak at about 44°C. The latter may correspond to the mixed complex indicated by the X-ray diffraction pattern. It should be noted that when the mole fraction of the salt n was 0.18 and 0.25, higher melting complexes were indicated by the DSC data for MEEP/PEO mixed electrolytes

containing LiBF_4 and LiClO_4 .⁽⁸⁾ The DSC data for 87 MEEP/13 PEGDA- $(\text{LiClO}_4)_{0.25}$ and 87 MEEP/13 PVP- $(\text{LiClO}_4)_{0.25}$ also confirmed their amorphous morphology. These data further indicated that the former is stable up to at least 200°C while the latter may decompose at about 190°C.

The structural data we have obtained suggest that generally speaking the crystalline-amorphous morphology of the mixed polymer electrolytes is related to that of the individual polymer electrolytes of which they are composed. $\text{LiN}(\text{SO}_2\text{CF}_3)_2$ complexes are an exception. The large imide anion appears to discourage precipitation of crystalline complexes so that the related electrolytes may be amorphous phases. The improved dimensional stability of the composite electrolytes compared to $\text{MEEP}-(\text{LiX})_n$ is attributable to the fact that the mechanically strong, second polymer phase dispersed into the film matrix reinforces its overall mechanical strength. The Li^+ seems to complex with MEEP as well as the second polymer phase in the mixed electrolyte which is in keeping with the fact that the electron donating heteroatom in both MEEP and the second polymer is oxygen bonded to carbon. The structural factors responsible for the unusual dimensional stability of $\text{MEEP}-(\text{LiAlCl}_4)_n$ are presently unknown. It is possible that the AlCl_4^- anions serve as crosslinks between the polymer chains, probably involving interactions with the $\text{P}=\text{N}$ bonds.

Ionic Conductivity and Transport: The specific conductivity versus temperature data presented in Figure 5 illustrate how the conductivity of a $\text{MEEP}/\text{Poly}(\text{olefin oxide})-(\text{LiX})_n$ mixed polymer electrolyte relates to that of electrolytes derived from the individual polymers. The conductivity versus $1/T$ behavior of $\text{MEEP}-(\text{LiBF}_4)_{0.25}$ and $\text{PPO}-(\text{LiBF}_4)_{0.13}$ is indicative of these polymers being fully amorphous. That is, the conductivity increases in a monotonic fashion with increasing temperature which, as previously discussed⁽¹⁴⁾, is in accordance with the Vogel-Tamann-Fulcher relationship, $\sigma = AT^{-1/2} \exp(-E_a/R(T-T_0))$ for

a predominantly amorphous polymer. There are two important features in the σ versus $1/T$ curves for PEO-(LiBF₄)_{0.125} and MEEP/PEO-(LiBF₄)_{0.125}: i) the cooling and heating curves show hysteresis; and ii) two distinct temperature regions appear in which the slopes of the curves which correspond to the activation energy for conduction have different values; below 60°C the slope is much steeper than above 60°C.

It is well known that the abrupt increase in conductivity at about 60°C in PEO-based electrolytes is due to the melting of crystalline PEO and the consequent formation of a uniformly plastic electrolyte phase. The term "plastic phase" is preferred here to indicate the more fluid nature of this phase, as opposed to the term "amorphous phase" which does not necessarily imply increased fluidity. Indeed, PPO-(LiBF₄)_{0.13} is fully amorphous over the same temperature of Figure 5, yet its conductivity at 75°C is significantly lower than that of the others. This is most probably due to a relatively lower amount of fluidity in this electrolyte. The usual argument based on poorer complexation of Li⁺ by PPO does not appear to be fully satisfactory in view of the rather similar conductivity of both PEO and PPO complexes below 60°C. It is also interesting that above 60°C, the conductivities of LiBF₄ complexes of PEO, PEO/MEEP mixture and MEEP are all rather similar. At this temperature, the three electrolytes appear to reach a rather similar state of "plasticity" whereby the large amplitude segmental motion of the polymer backbone which assists the conductivity of the ions becomes appreciable. As with PEO-(LiX)_n, MEEP/PEO-(LiX)_n electrolytes also exhibit hysteresis in their σ vs $1/T$ plots associated with heating and cooling. During cooling, the shape of the plot was altered significantly with almost a complete elimination of the knee between 50 and 60°C. The heated electrolyte was found to maintain its higher plasticity for an extended period of time. The original morphology return when the electrolyte was stored at room temperature for about a month.

The activation energy for conduction in MEEP/PEO-(LiBF₄)_n was determined as a function of the ratio of the two polymers and the salt concentration (8). For the 55/45 polymer weight ratio, the activation energy below 60°C was insensitive to salt concentrations corresponding to n between 0.016 and 0.18. However, activation energy increased substantially when the amount of MEEP decreased from 70 w/o to 50 w/o. This behavior is consistent with higher plasticity for electrolytes having higher MEEP contents. Above 60°C, correlations between activation energy and electrolyte composition were not straightforward.

The complex impedance spectra of electrolytes using stainless steel blocking electrodes appear to reflect the degree of plasticity of the electrolytes. Conductivities of the electrolytes were calculated from such spectra. Figure 6a and 6b display the data for 55 MEEP/45 PEO-(LiBF₄)_{0.13} at different temperatures. The AC frequency range was between 6 Hz and 100 kHz. For fresh electrolytes at room temperature, the Cole-Cole spectrum represented a semicircle with a spur at low frequencies. This represents an equivalent circuit in which the electrode capacitance is in series with a parallel combination of electrolyte resistance and its capacitance. With increasing temperature the semicircle gave way to the spur and at a temperature above the melting point of PEO only a straight line usually remained. At these temperatures, the capacitive component of the electrolyte appears to be negligible up to a frequency of 100 kHz. The high plasticity induced at temperatures above the melting point of PEO appears to decrease the dielectric relaxation times of the electrolyte so that in our frequency range of impedance measurement the electrolyte capacitance becomes negligible. In fact, as shown in Figure 6a, the electrolyte heated once to 80°C and then cooled to 20°C shows an impedance spectrum indicative

of a highly plastic electrolyte. Similarly, a high degree of plasticity of MEEP-(LiX)_n as the cause of their high room temperature conductivity is reflected in their impedance spectra, an example of which is given in Figure 7.

It can be seen from the data in Figure 8 that the conductivities of the other mixed electrolytes MEEP/PEO-(LiX)_n, MEEP/PVP-(LiX)_n, MEEP/PEGDA-(LiX)_n are also very high at room temperature, being only about an order of magnitude lower than that of MEEP-(LiX)_n. Conductivities at 25 and 50°C for a group of electrolytes are listed in Table 2. The data illustrate some interesting effects of Li salt on conductivity. The dimensionally stable films of 55 MEEP/45 PEO-[LiN(SO₂CF₃)₂]_{0.13} exhibited a room temperature conductivity higher than that of all the MEEP-(LiX)_n complexes except MEEP-[Li(SO₂CF₃)₂]_{0.13} whose conductivity is identical to that of the MEEP/PEO mixed electrolyte containing the imide salt. The complex impedance spectrum of 55 MEEP/45 PEO-(LiN(SO₃CF₃)₂)_{0.13} had a profile similar to that shown in Figure 7 indicative of a highly plastic material.

The conductivities of LiAlCl₄ complexes of MEEP are presented in Figure 9 as plots of σ vs $1/T$. Despite its good mechanical strength, the conductivity of MEEP-(LiAlCl₄)_{0.13} at room temperature is almost the same as that of MEEP-(LiClO₄)_{0.25}. The corresponding impedance spectra in Figure 10 seems to correlate this high conductivity with the high degree of plasticity of the electrolyte. At higher LiAlCl₄ concentrations the conductivity decreases perhaps due to increased ion-pairing.

The results we have obtained suggest that the dimensionally stable electrolytes derived from LiN(SO₂CF₃)₂ and LiAlCl₄ retain good plasticity even in a solid matrix to produce high conductivity at room temperature in the corresponding electrolyte films. Evidently, polymer electrolytes having an ability to retain a high degree of plasticity or microfluid domains in a solid matrix are the ones expected to be highly ionically conductive

while being dimensionally stable. Indeed, the higher room temperature conductivity of the MEEP/PVP-(LiClO₄)_{0.25} composite electrolyte compared to its PEO or PPO composite counterpart may be ascribed to a predominantly plastic phase of MEEP-(LiClO₄)_n contained in a solid matrix in which PVP primarily is a matrix reinforcement to impart mechanical strength to the electrolyte.

The results presented above demonstrate that several convenient methods are now available for the preparation of dimensionally stable, MEEP-based, free standing electrolytes whose conductivities are comparable to that of MEEP-(LiBF₄)_{0.25} and MEEP-(LiClO₄)_{0.25} which have poor mechanical stability. The good mechanical strength of the LiAlCl₄ complexes is interesting, especially in view of the fact that they did not exhibit any significant amount of crystallinity. It may be noted here that the behavior of MEEP/PEO mixed polymer electrolytes containing LiAlCl₄ we reported previously⁽⁸⁾ was somewhat peculiar. For example, the conductivity of MEEP/PEO-(LiX)_{0.13} electrolytes decreased in the order LiBF₄ ≈ LiClO₄ > LiCF₃SO₃ > LiAsF₆ > LiAlCl₄. The LiAlCl₄-containing films were all characterized by more rigidity than the others, probably indicating the unusual interaction of the AlCl₄⁻ anion with the polymer chain.

The Li⁺ transport number (*t_s*) of some selected MEEP-based mixed polymer electrolytes was determined using the potentiostatic polarization technique [11]. Table 3 lists the values for electrolytes having different polymer compositions and two different Li salts, namely LiClO₄ and LiBF₄. As can be seen in this table, transport numbers around 0.40 were obtained for Li⁺ in these electrolytes. The reason for the somewhat higher transport number of the LiBF₄ complex of MEEP/PEO is not clear at this time. Previously,⁽¹⁴⁾ *t_s* of 0.32 and 0.17, respectively, were obtained for MEEP-(LiCF₃SO₃)_{0.17} and MEEP-(LiBF₄)_{0.17} electrolytes. It is worth noting also that the transport numbers in PEO-(LiX)_n^(16,17) electrolytes

ranged between 0.4 and 0.5. Since the mobile cations in polymer electrolytes are triple and/or quadruple ions, the t_+ reported here does not strictly represent transport number for Li^+ . It is more accurately described as the DC conductivity which transports Li ions from the negative to the positive electrode.

The higher the ionic conductivity for an electrolyte the lower the resistive polarization (iR drop) during discharge of a Li cell utilizing the electrolyte. On the other hand, a transport number of less than one for Li^+ implies migration of the anion to the Li electrode during cell discharge until a steady state discharge current is reached. This results in the accumulation of Li salt at the Li-electrolyte interface and possible interfacial polarization of the Li electrode. The effect of Li^+ transport number on the steady state limiting current of a Li cell, e.g., a Li/TiS_2 cell, can be described in terms of the relationship in equation [1]⁽¹⁷⁾. In this equation, D_{Li}^+ , C_b , and t_+ are

$$I_l = \frac{2nFD_{\text{Li}}^+C_b}{d(1-t_+)} \quad (1)$$

the diffusion coefficient, the bulk concentration and transport number of Li^+ , and d is the interelectrode spacing which includes the thickness of the porous cathode. An optimized solid-state cell will utilize a 0.05 mm thick electrolyte film and a 0.1 mm thick cathode. The concentration of Li^+ in the 55 MEEP/45 PEO- $(\text{LiClO}_4)_{0.13}$ electrolyte is 1.8 moles/liter. Using a diffusion coefficient of $10^{-7} \text{ cm}^2/\text{sec}$ for Li^+ at 70°C and a Li^+ transport number of 0.4, the steady-state limiting current can be calculated to be 0.6 mA/cm^2 . Equation 1 shows that the limiting current can be increased by adjusting a variety of parameters including a higher salt concentration, and higher diffusion coefficient and transport number for Li^+ .

Electrochemical Stability: Cyclic voltammetric studies at stainless steel or gold working electrodes were carried out to assess the electrochemical stability domain of the

MEEP-based mixed polymer electrolytes. Ideally, a polymer electrolyte should exhibit a stability domain between 0.0 and 4.0V, the potential region where the electrochemical reactions of the Li negative and the positive electrode occur. Figure 11a shows the voltammetric scan of a 55 w/o MEEP:45 w/o PEO-(LiClO₄)_{0.13} at a scan rate of 2 mV/s at 75°C. The electrode was stainless steel. The electrode was swept first cathodically from the open-circuit potential of 2.8V to -1V, followed by an anodic sweep up to 5V. The potentials reported here are with respect to the Li⁺/Li electrode and are without iR compensation.

The first small rise in reduction current took place at about 1.4V, followed by another small cathodic current peak beginning at 0.7V. Both of these peaks appear to be associated with corresponding anodic peaks of much smaller magnitude at about 2.2 and 1.3V, respectively. We have observed peaks at potentials very close to these also in the cyclic voltammograms of MEEP/PPO-LiClO₄⁽⁹⁾, polytetrahydrofuran-LiClO₄⁽¹⁸⁾ and poly(1,3-dioxolane)-LiClO₄ electrolytes⁽¹⁸⁾. The common specie in these electrolytes is LiClO₄, suggesting that these peaks may be associated with the reduction and oxidation of LiClO₄. It appears that the polymer itself undergoes little reduction at these low potentials. The sharp rise in cathodic current beginning at -0.2V is due to the reduction of Li⁺ and the deposition (plating) of the resulting Li on the stainless electrode, equation [2].



The anodic peak centered around 1.5V can be ascribed to the stripping (oxidation) of the Li. Comparison of the quantities of charge involved in Li plating and stripping reveals that the processes are fully chemically reversible. A very small anodic peak around 4.5V was observed on further scanning the electrode up to a potential of 5.0V. As shown in Figure 11b, voltammograms similar to these were also obtained with the MEEP/PEGDA-LiClO₄ electrolyte. The effect of Li salt on electrolyte stability was investigated with the cyclic

voltammogram of 55 MEEP/45 PEO-(LiBF₄)_{0.13} (Figs. 12a and 12b). The Li stripping process was followed by another closely lying peak which assumed a distinct shape on continued cycling (Fig. 12b). Despite the two steps associated with the stripping process, Li cycling appeared to be fully chemically reversible. We can explain the two steps in the Li stripping process as follows. Li being plated on the stainless steel electrode is in two different environments: a first layer adjacent to the stainless steel electrode and strongly adsorbed to it; a second fraction residing on the first layer of Li. Consequently, the second fraction of Li is stripped at a potential closer to the Li⁺/Li potential than the first fraction which is stripped at a slightly more positive potential.

Figure 13 presents the cyclic voltammogram of 55 MEEP/45 PEO-(LiClO₄)_{0.13} electrolyte at a gold working electrode. Again, as in Figure 11, there are two very small peaks around 1.5 and 0.9V prior to the main reduction peak beginning at around 0.2V. The oxidation peak corresponding to the reduction at 0.2V was observed around 0.5V, and the reduction and oxidation processes appeared to be chemically reversible. Unlike on a stainless steel electrode where the plating of Li occurred, as expected, at a potential negative of Li⁺/Li, Li plating on Au takes place at a potential positive of Li⁺/Li. This behavior is similar to that found when the electrode is Al or other metals which can alloy with Li⁽¹⁹⁾. Indeed, Li and Au forms an alloy of the composition Li₃Au⁽¹⁹⁾. The alloying of Li with and dealloying of Li from Au at 75°C exhibits good chemical reversibility. On further scanning the electrode anodically, an appreciable current began to appear at around 4.7V. On a return sweep, there was a cross-over of the oxidation and the reduction peaks, presumably due to electrode passivation from adsorption of electroactive species on the electrode. The gold electrode, unlike stainless-steel, appeared black after excursion to the Li plating potential, supporting the formation of an alloy of Li and Au.

The cyclic voltammogram of the 55 MEEP/45 PEO-(LiBF₄)_{0.07} electrolyte on a glassy carbon electrode showed a reduction peak at 2.1V, followed by two small reduction peaks at 1.35 and 0.45V. The current involved in these reduction processes on carbon was higher than that observed on the metal electrodes and may be indicative of a catalytic effect of carbon. A peak at 0.45V, probably corresponding to the intercalation of Li into C was also observed. The latter process is typically observed in the voltammograms of carbon in non-aqueous electrolytic solutions containing Li salts. The practical relevance of the electrochemistry of polymer electrolytes on a carbon surface lies in the fact that high surface area carbon is used as an additive in the cathode structure of Li batteries to enhance electronic conductivity to the electrode. Any redox processes involving the electrolyte on the cathode can affect battery performance. The anodic stability of the electrolytes on carbon exceeded 4.25V versus Li⁺/Li.

The cycling of the 55 MEEP/45 PEO-(LiBF₄)_{0.13} electrolyte on a Li metal substrate is illustrated in Figure 14. The plating and stripping of Li are the only electrochemical processes occurring here. Practically no reduction of the electrolyte is observed. This is extremely significant since unwanted side reactions of the electrolyte which can interfere with Li plating and stripping will be of little concern here.

Solid-State Li/TiS₂ Cells: Li/TiS₂ cells utilizing the dimensionally stabilized MEEP-electrolytes were fabricated, and galvanostatically discharged and charged (cycled) at several temperatures including 20, 50 and 75°C. Since MEEP-(LiX)_n itself is dimensionally unstable, it was immobilized by incorporating into the pores of a fiberglass filter paper⁽⁴⁾. This enabled comparisons of the rate capabilities of neat MEEP electrolytes with MEEP-based composite electrolytes.

The open-circuit potentials of freshly prepared cells were between 2.8 and 2.9V, which on heating to 50°C declined slightly to values between 2.6 and 2.8V. Figure 15 shows typical constant current charge-discharge curves of a Li/TiS₂ cell using MEEP-(LiClO₄)_{0.25} electrolyte supported in a fiber-glass separator paper. The cells were discharged to a voltage limit of 1.6V and charged to 3.0V. The data encompass three different rates at 50°C. The current densities were 0.05, 0.1 and 0.2 mA/cm² at the C/28, C/15 and C/7 rates. The cell delivered the capacities of 1.10, 0.45 and 0.28 mAh/cm², respectively, at these rates. Increasing the discharge temperature from 50 to 75°C more than doubled the capacity of the cell. The mid-discharge voltage was 2.1V, while the average charging potential was 2.5V. The coulombic efficiency during the cycling was between 90 and 100%.

Figure 16 shows the long-term cycling performance of a second Li/TiS₂ cell using MEEP-(LiClO₄)_{0.25} supported on a fiber-glass paper matrix. The cell was cycled more than 200 times. The continuous decrease in capacity with cycling is somewhat typical of unoptimized laboratory cells in which the capacity loss is associated with several factors including cathode structural problems, inefficiencies in the recharging of the anode and cathode, and any number of mechanical reasons. However, the present data clearly show that the MEEP-based electrolytes are chemically and electrochemically stable in contact with Li and can be fabricated into long cycle life secondary Li batteries.

A knowledge of the overcharge and overdischarge behavior of a secondary Li cell, that is, discharge and charge into potential regions beyond where the normal electric current producing redox chemistry of the cell occurs, is important since such a condition could be encountered in the normal operation of a battery. Undesirable electrochemical reactions during overdischarge and overcharge can be detrimental to the stability of the cell. Figure 17 depicts that the cell can function normally even after a considerable extent of overcharge.

In fact, following the overcharge which occurred inadvertently at the 76th cycle due to the malfunction of the cycling equipment, the cell had been discharged down to 0.0V before being charged back in the 77th cycle. The overdischarge, as well as the overcharge, did not have a deleterious effect on subsequent cell performance.

The performance of a Li/TiS₂ cell using a mixed polymer electrolyte is illustrated with the MEEP-PVP-(LiClO₄)_{0.25} electrolyte, at three different temperatures and 0.05 mA/cm² (C/20) (Fig. 18). At room temperature the cell delivered about 17% (0.15 mAh/cm²) of its theoretical cathode capacity of 0.88 mAh/cm². At 75°C, the cathode utilization was 100%. The charging efficiency at room temperature was 85%, while at the higher temperature it was 100%, indicating slower recharge kinetics at low temperatures. The rate capability of the cell containing MEEP/PVP-(LiClO₄)_{0.25} was very much similar to that of the cell described above utilizing MEEP-(LiClO₄)_{0.25}.

Cells utilizing MEEP/PEO-(LiClO₄)_{0.13} composite electrolytes exhibited rate capabilities inferior to those described above. For example, a Li/TiS₂ cell utilizing 70 w/o MEEP:30 w/o PEO-(LiClO₄)_{0.12} could not be discharged at room temperature at a practical rate such as C/20 or C/40. However, when discharged at 65°C, about 86% of the theoretical cathode capacity, i.e., 1.2 mAh/cm² was obtained at the C/40 rate, while at 50°C the utilization was 0.96 mAh/cm². The coulombic efficiency during cycling was close to 100%.

CONCLUSIONS

Several, relatively simple methods have been developed to prepare dimensionally stable MEEP-based electrolytes. Pure MEEP when complexed with LiAlCl_4 forms dimensionally stable, free-standing thin films. This contrasts the behavior of MEEP complexes we have prepared with many other commonly used Li salts. Additional methods of dimensional stabilization include blending $\text{MEEP}^{2+}(\text{LiX})_n$ with a poly(olefin oxide) such as PEO or PPO, or forming composites with photo-polymers such as PEGDA and PVP. Some of these electrolytes exhibited conductivities close to those of the dimensionally unstable $\text{MEEP}^{2+}(\text{LiX})_n$ indicating that they retain a high degree of plasticity or domains of microfluidity. Transport numbers ranging between 0.4 and 0.5 have been determined for Li^+ in these electrolytes which are very similar to that obtained in other Li^+ -conductive polymer electrolytes. The dimensionally stabilized MEEP-based electrolytes have demonstrated good electrochemical stability on a number of metal substrates including Li. The long-term cycling capability of Li/TiS_2 secondary solid-state cells fabricated with these electrolytes indicate that they are useful for low to moderate rate applications at ambient temperature.

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TABLE 1
X-RAY DIFFRACTION DATA FOR LiClO_4 , PEO AND POLYMER ELECTROLYTES

LiClO_4 , d (Å)	PEO, d (Å)	PEO-(LiClO_4) _{0.13} , d (Å)	MEEP/PEO-(LiClO_4) _{0.13} , d (Å)			
			40/60 ³	50/50	60/40	70/30
		3.00 ₁₀₀	3.02 ₆₀	3.02 ₇₀	3.01 ₈₀	3.03 ₁₀₀
	2.31 ₇₀	2.32 ₄₀				
2.16 ₂₀						
			2.02 ₁₀	2.04 ₆₀	2.02 ₄₀	2.02 ₉₀
	1.92 ₁₀₀	1.93 ₈₀	1.96 ₁₀₀	1.96 ₁₀₀	1.96 ₁₀₀	1.96 ₉₀
1.82 ₁₀₀	1.80 ₂₀					
1.76 ₃₀	1.73 ₁₀					
1.40 ₂₀				1.60 ₁₀	1.60 ₁₀	
1.30 ₂₀						
1.24 ₂₀					1.48 ₆	
1.23 ₃₀						
1.09 ₁₀						

¹The subscript refers to the relative intensity of the X-ray lines.

²All the electrolytes were X-rayed at least four weeks after they were prepared.

³Weight ratio of polymers in percentage.

TABLE 2

CONDUCTIVITIES OF A NUMBER OF MEEP-BASED ELECTROLYTES
AT 25 AND 50°C

Polymer Electrolytes	Conductivity, $\Omega^{-1}\text{cm}^{-1}$	
	25°C	50°C
MEEP-(LiClO ₄) _{0.25}	1.7×10^{-5}	5.0×10^{-5}
55 w/o MEEP/45 w/o PEO-(LiClO ₄) _{0.13}	1.3×10^{-6}	9.0×10^{-6}
55 w/o MEEP/45 w/o PPO-(LiClO ₄) _{0.13}	9.0×10^{-8}	3.5×10^{-6}
MEEP-LiN(CF ₃ SO ₂) ₂	6.5×10^{-5}	1.6×10^{-4}
55 w/o MEEP/45 w/o PEO- [LiN(CF ₃ SO ₂) ₂] _{0.13}	6.7×10^{-5}	1.2×10^{-4}
87 w/o MEEP/13 w/o PVP-(LiClO ₄) _{0.13}	4.0×10^{-6}	2.3×10^{-5}
87 w/o MEEP/13 w/o PEGDA-(LiClO ₄) _{0.13}	1.2×10^{-6}	7.2×10^{-6}
MEEP-(LiAlCl ₄) _{0.13}	1.2×10^{-5}	4.0×10^{-5}

TABLE 3

TRANSPORT NUMBERS FOR Li^+ IN SOME MEEP-BASED
ELECTROLYTES

Cell Number	Electrolyte	t_+
1	87/13 MEEP/PEGDA- $(\text{LiClO}_4)_{0.13}$	0.40
2	70/30 MEEP/PEO- $(\text{LiClO}_4)_{0.13}$	0.44
3	70/30 MEEP/PEO- $(\text{LiClO}_4)_{0.13}$	0.36
4	70/30 MEEP/PEO- $(\text{LiBF}_4)_{0.13}$	0.40
5	55/45 MEEP/PEO- $(\text{LiBF}_4)_{0.13}$	0.39
6	55/45 MEEP/PEO- $(\text{LiBF}_4)_{0.13}$	0.34
7	55/45 MEEP/PPO- $(\text{LiBF}_4)_{0.13}$	0.52
8	55/45 MEEP/PPO- $(\text{LiClO}_4)_{0.13}$	0.40

FIGURE CAPTIONS

- Figure 1. Comparison of the IR spectra of PEO and PEO-(LiClO₄)_{0.13}.
- Figure 2. IR spectra of MEEP, MEEP-(LiClO₄)_{0.25} and MEEP-(LiAlCl₄)_{0.25}.
- Figure 3. DSC traces of MEEP-(LiAlCl₄)_n complexes.
- Figure 4. DSC traces of PEO-(LiClO₄)_{0.125} and 55 MEEP/45 PEO-(LiClO₄)_{0.13}.
- Figure 5. Conductivities of the LiBF₄ complexes of MEEP, PEO, PPO and a MEEP/PPO mixture.
- Figure 6. (a) Room temperature complex impedance spectra for the electrolyte 55 MEEP/45 PEO(LiBF₄)_{0.13}, recorded between stainless steel blocking electrodes as fresh and after heating to 80°C followed by cooling to room temperature. (b) fresh electrolyte recorded at 45, 55 and 65°C.
- Figure 7. Room temperature complex impedance spectrum of MEEP-(LiClO₄)_{0.25} recorded between 6 Hz and 100 kHz.
- Figure 8. A comparison of the conductivities of several MEEP-based electrolytes.
- Figure 9. Conductivities of MEEP-(LiAlCl₄)_n complexes.
- Figure 10. Complex impedance spectra at room temperature for LiAlCl₄ complexes of MEEP, between 6 Hz and 100 kHz.
- Figure 11. Cyclic voltammogram of a) 55 MEEP/45 PEO-(LiClO₄)_{0.13} and b) 87 MEEP/13 PEGDA-(LiClO₄)_{0.13} on a stainless steel electrode at 75°C. Scan rate = 2 mV/s.
- Figure 12. Cyclic voltammogram of 55 MEEP/45 PEO-(LiBF₄)_{0.13} on a stainless steel electrode at 75°C. Scan rate = 2 mV/s. (a) first scan, (b) several scans beginning with the second scan.

- Figure 13. Cyclic voltammogram of 55 MEEP/45 PEO-(LiClO₄)_{0.13} at a gold electrode at 75°C. Scan rate = 0.5 mV/s.
- Figure 14. Cyclic voltammogram of 55 MEEP/45 PEO-(LiBF₄)_{0.13} on a Li electrode at 70°C. Scan rate = 2 mV/s.
- Figure 15. Capacity-rate behavior of a Li/TiS₂ cell using fiberglass matrix supported MEEP-(LiClO₄)_{0.25} electrolyte at 50°C.
- Figure 16. Long-term cycling behavior of a Li/TiS₂ cell using fiberglass matrix supported MEEP-(LiClO₄)_{0.25} electrolyte at 50°C. Current = 0.2 mA/cm², i_c = 0.05 mA/cm². Voltage limits = 1.6-3.0V.
- Figure 17. The effect of overcharge on the cycling of a Li/TiS₂ cell utilizing MEEP-(LiClO₄)_{0.25}. Current is same as in Figure 16.
- Figure 18. Constant current discharges of a Li/TiS₂ cell using 87 MEEP/13 PVP-(LiClO₄)_{0.25} electrolyte at C/20 and different temperatures. i_d = 0.08 mA/cm², i_c = 0.04 mA/cm². Voltage limits = 1.6-3.0V.

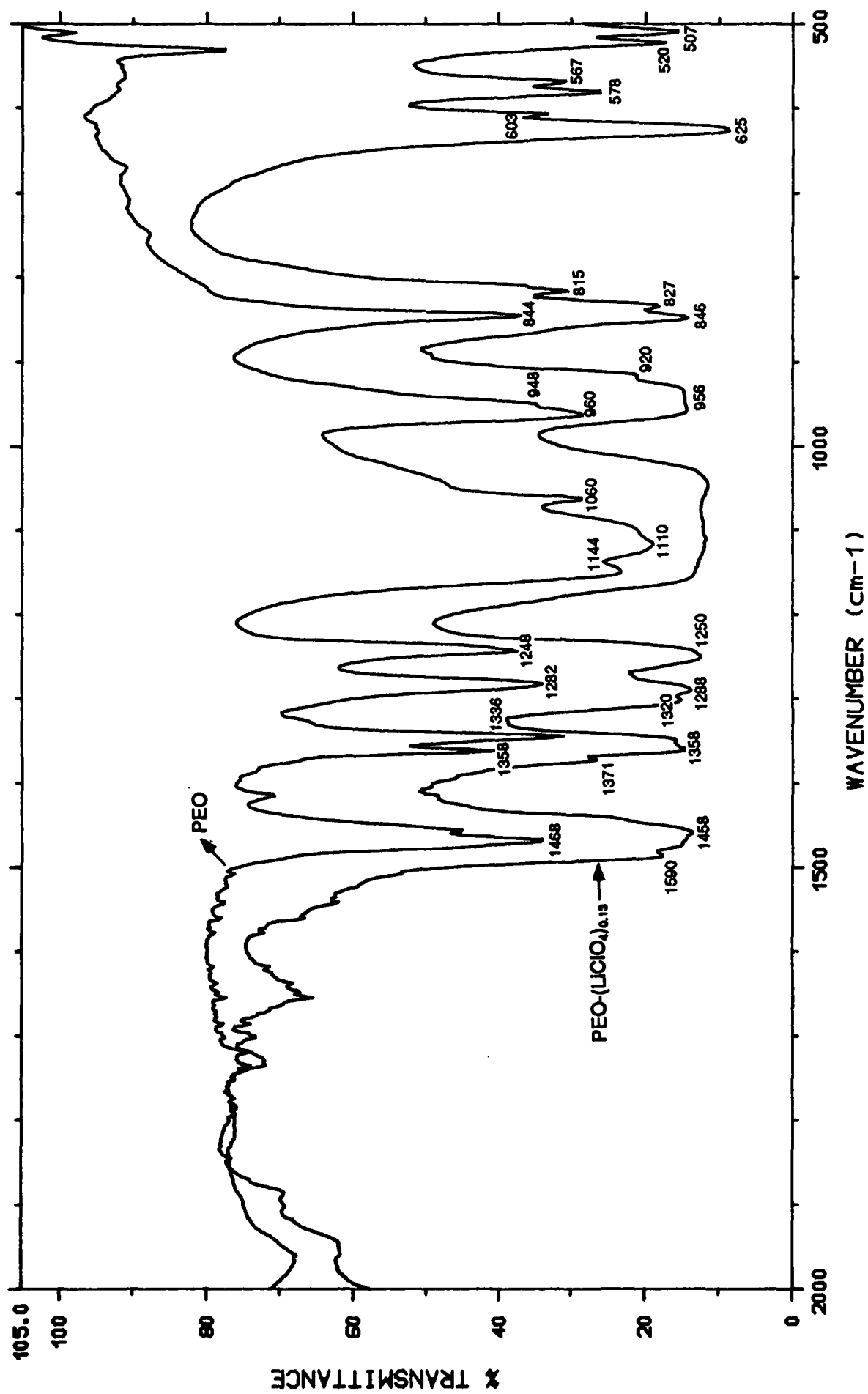


FIGURE 1

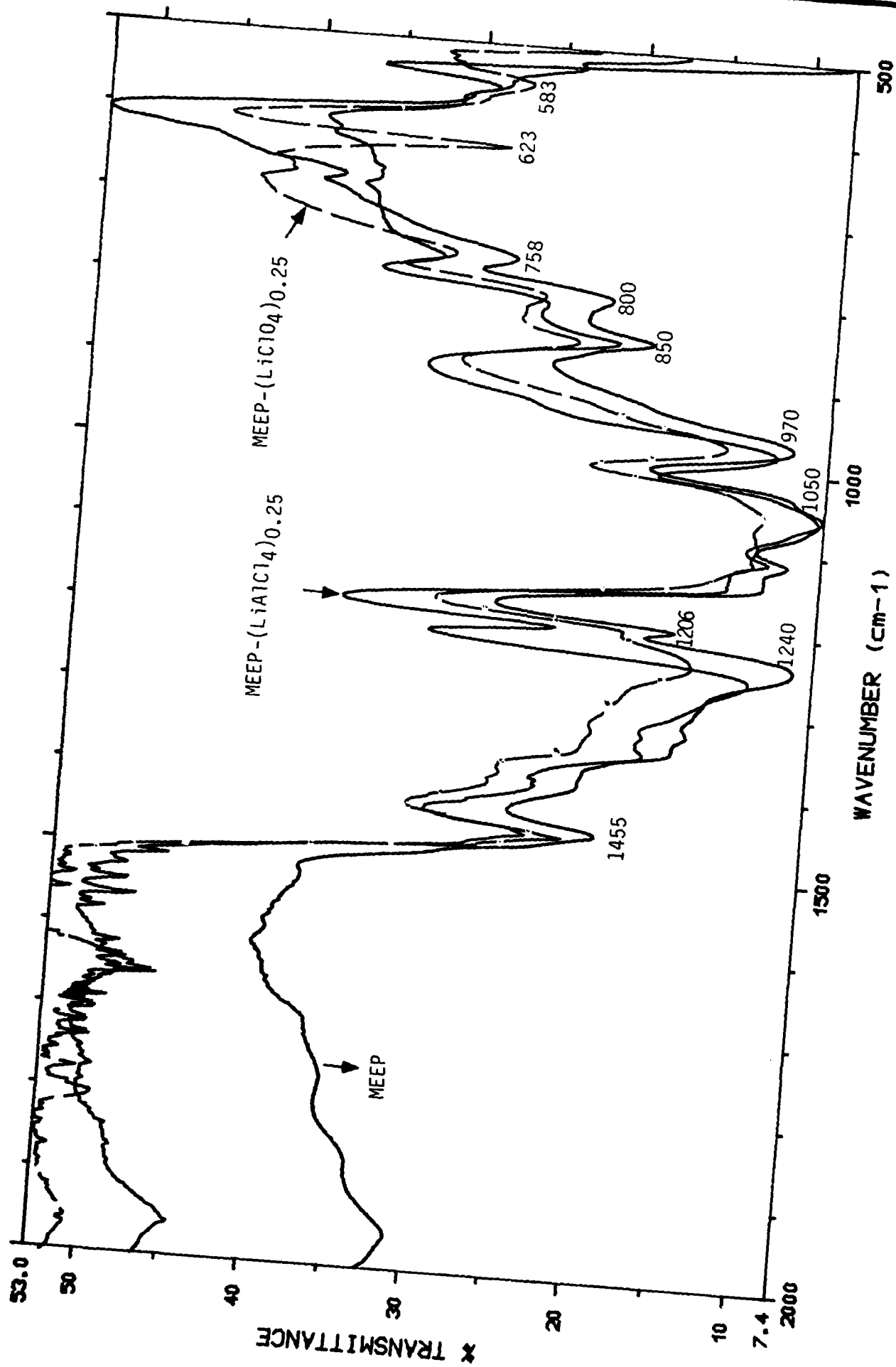


FIGURE 2

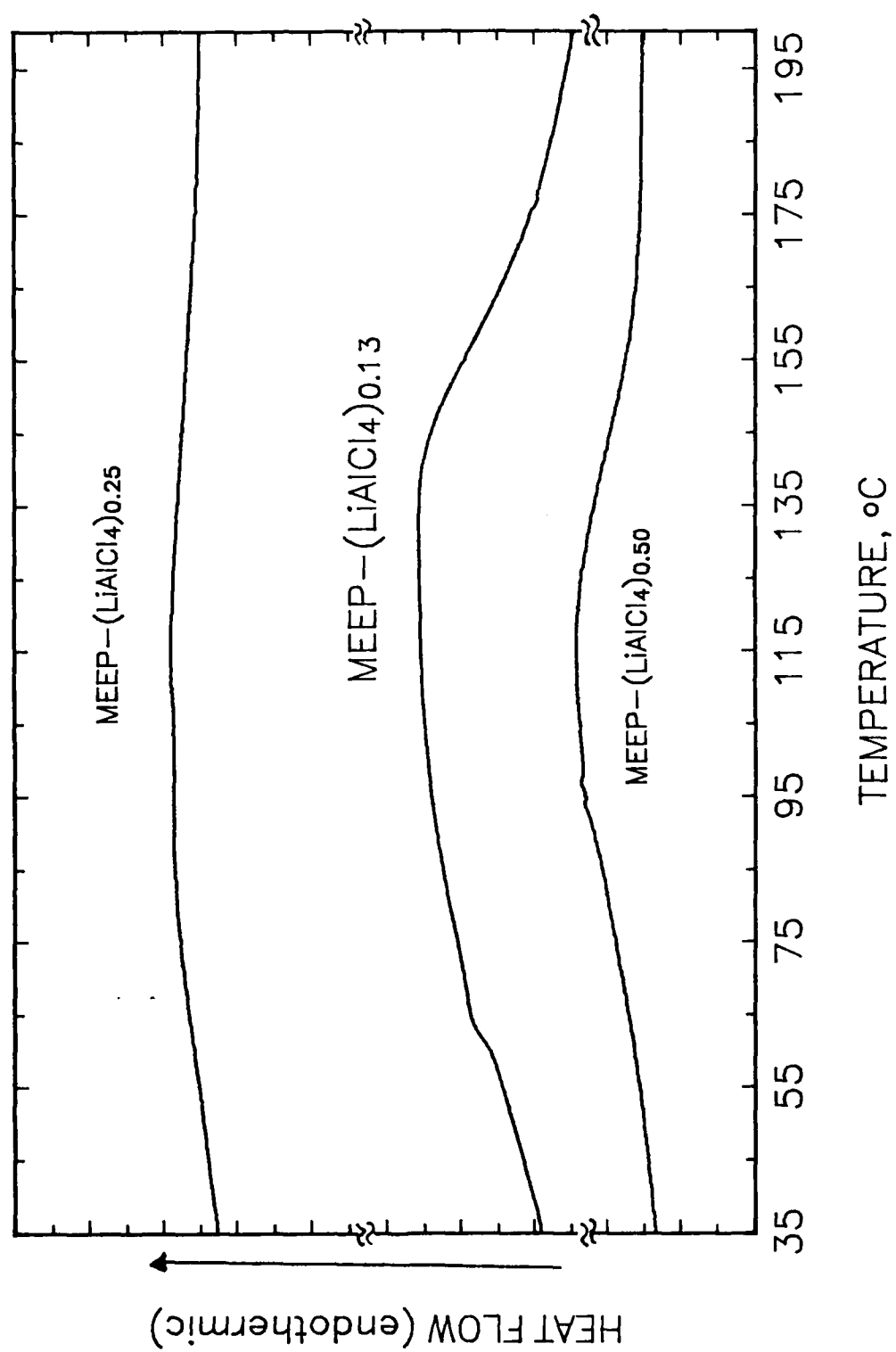


FIGURE 3

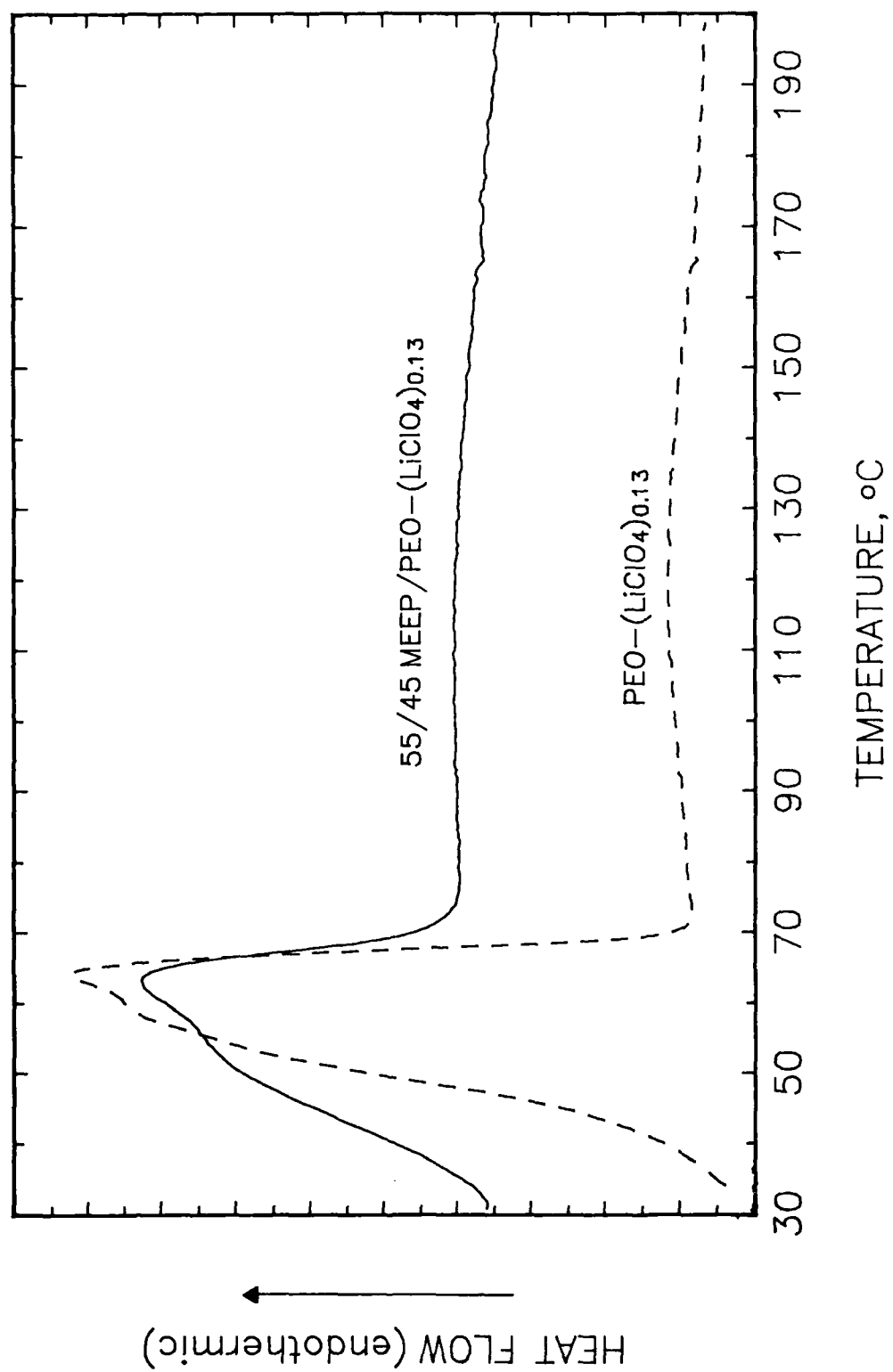


FIGURE 4

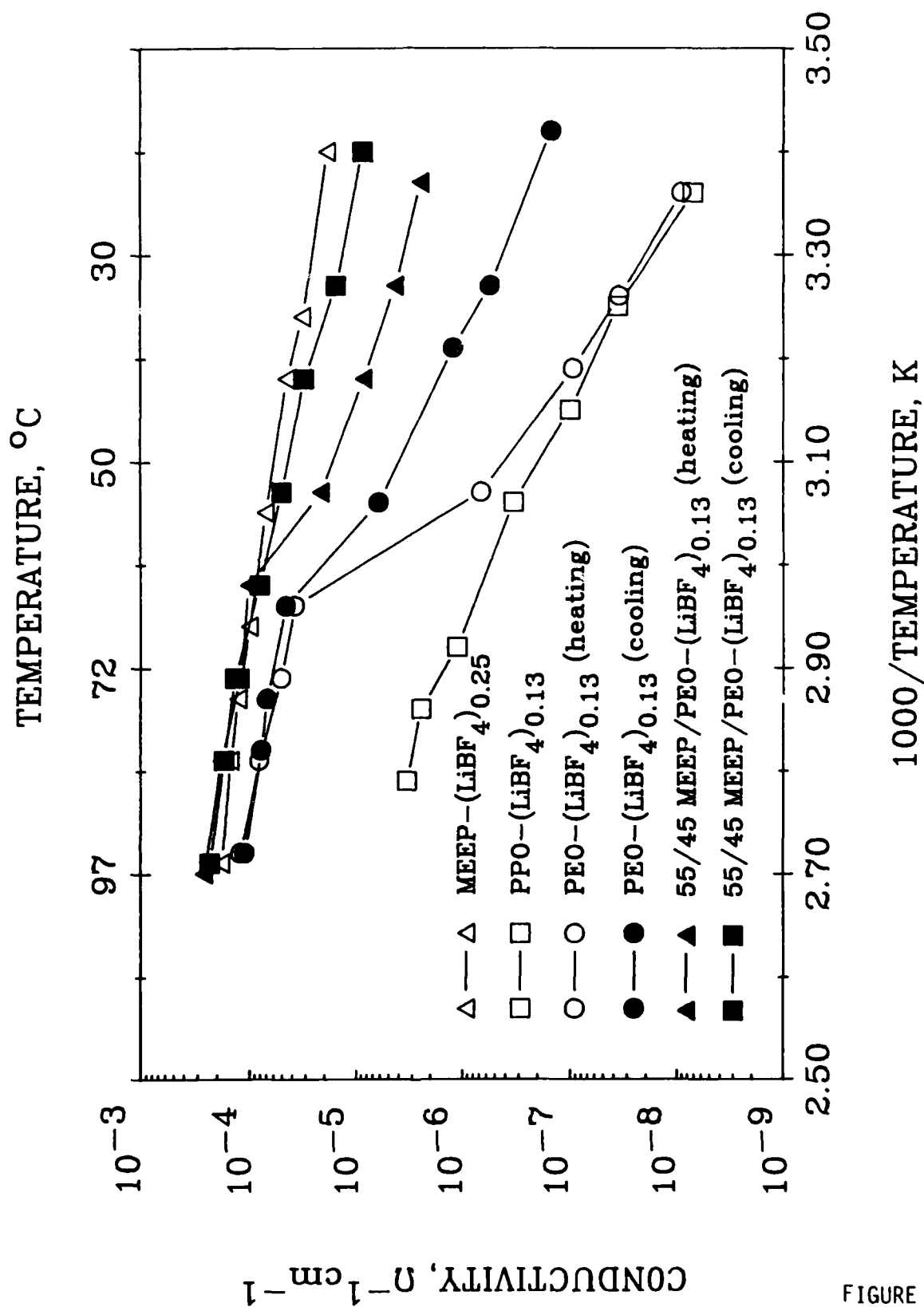


FIGURE 5

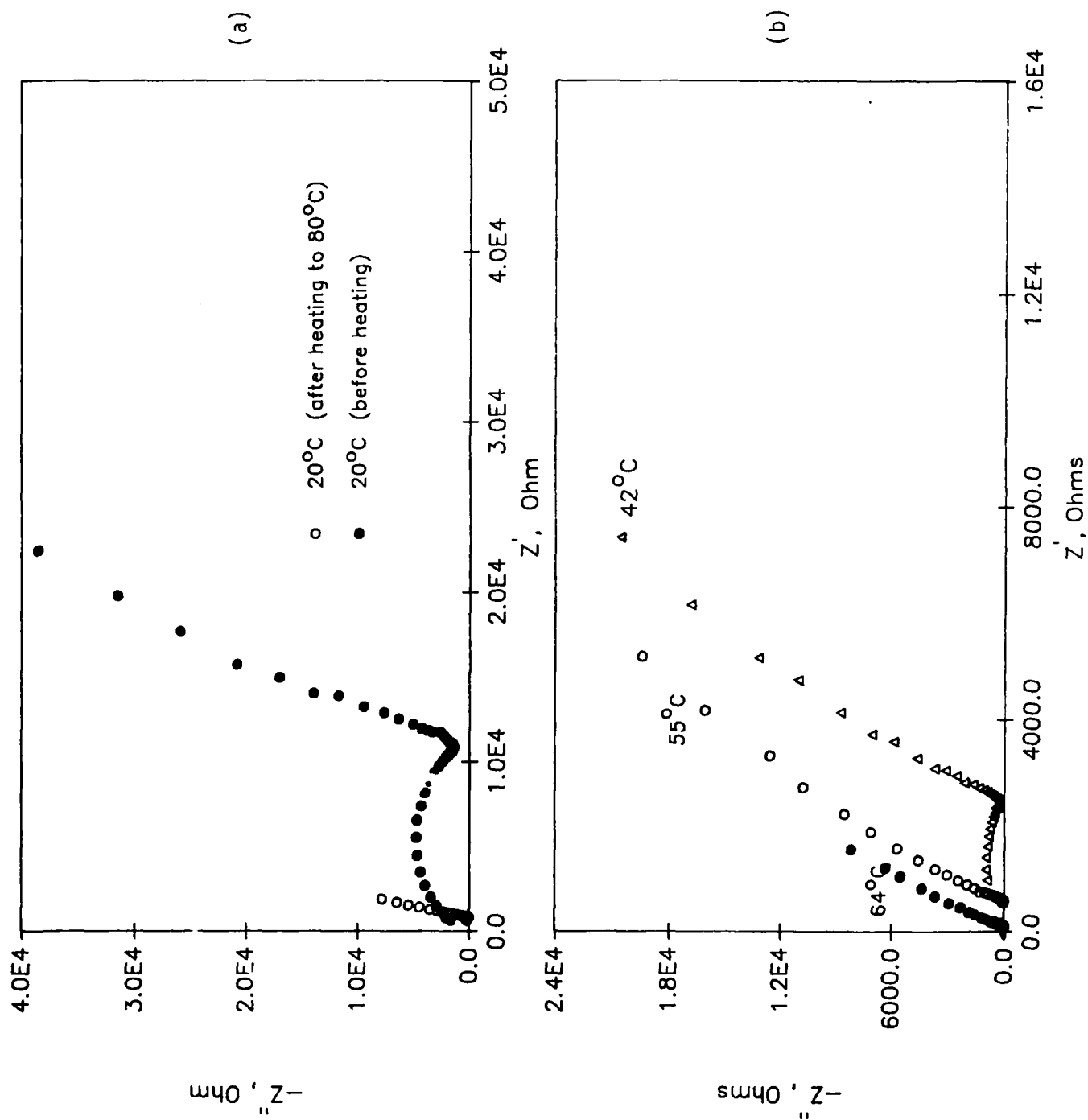


FIGURE 6

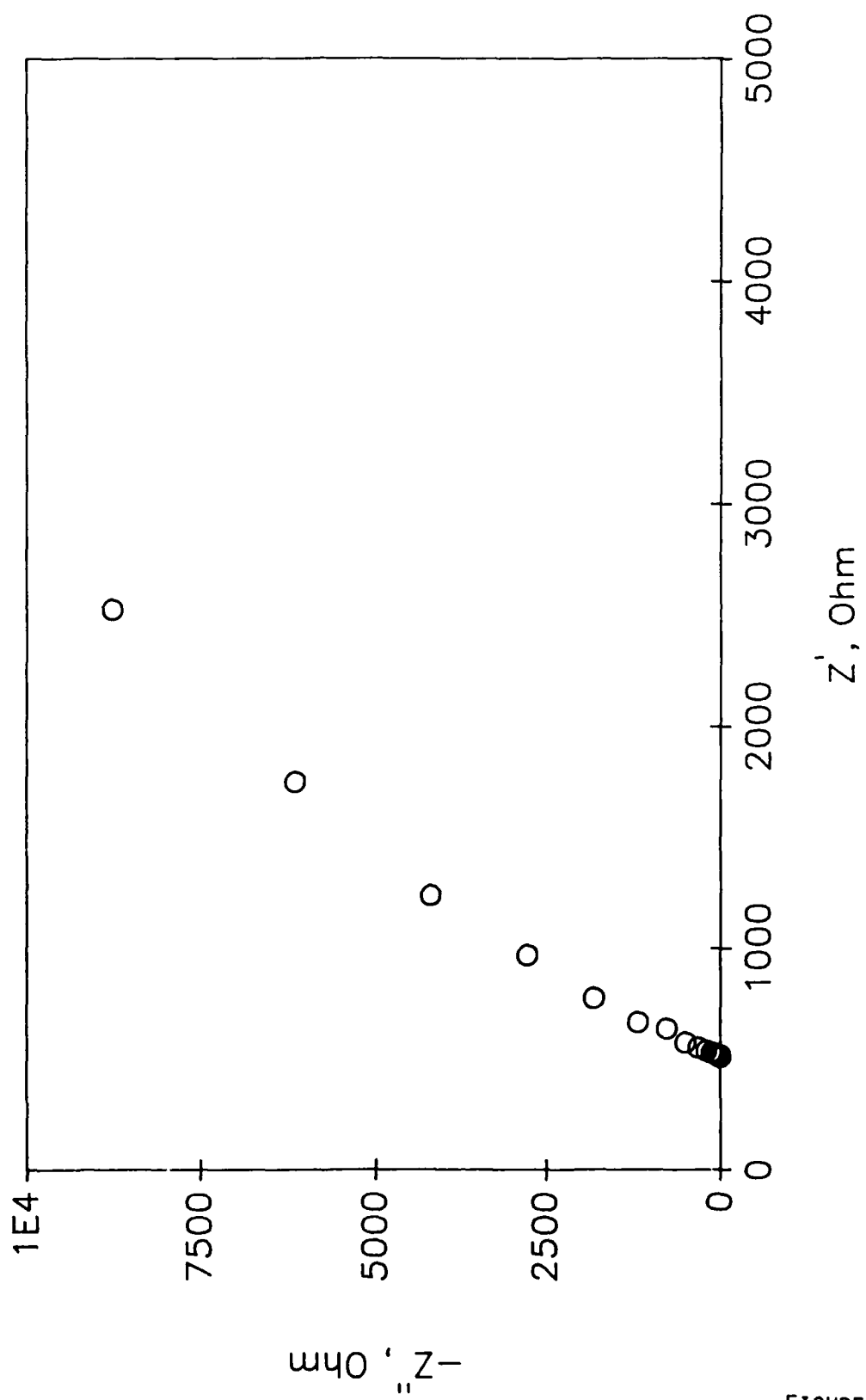


FIGURE 7

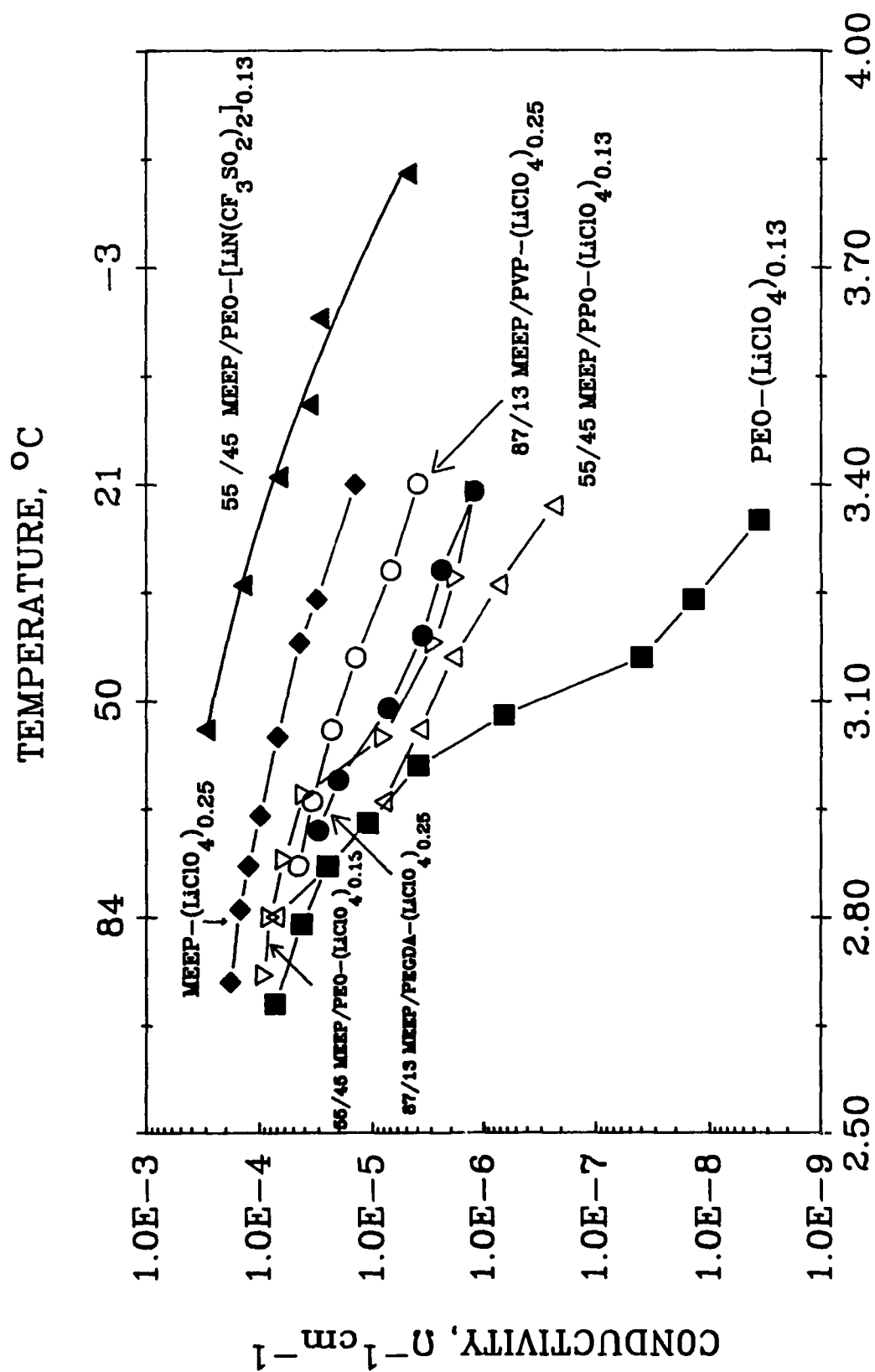


FIGURE 8

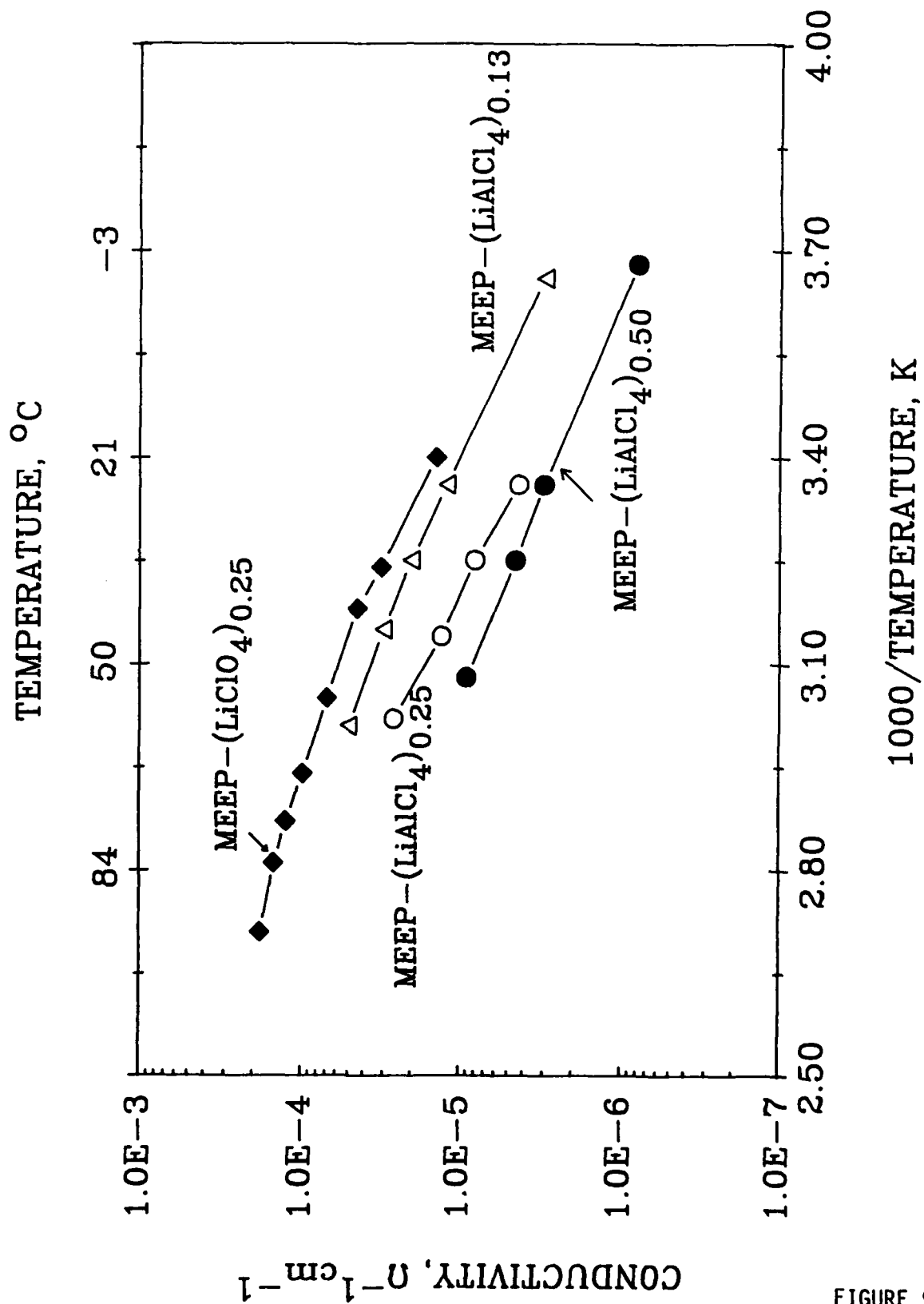


FIGURE 9

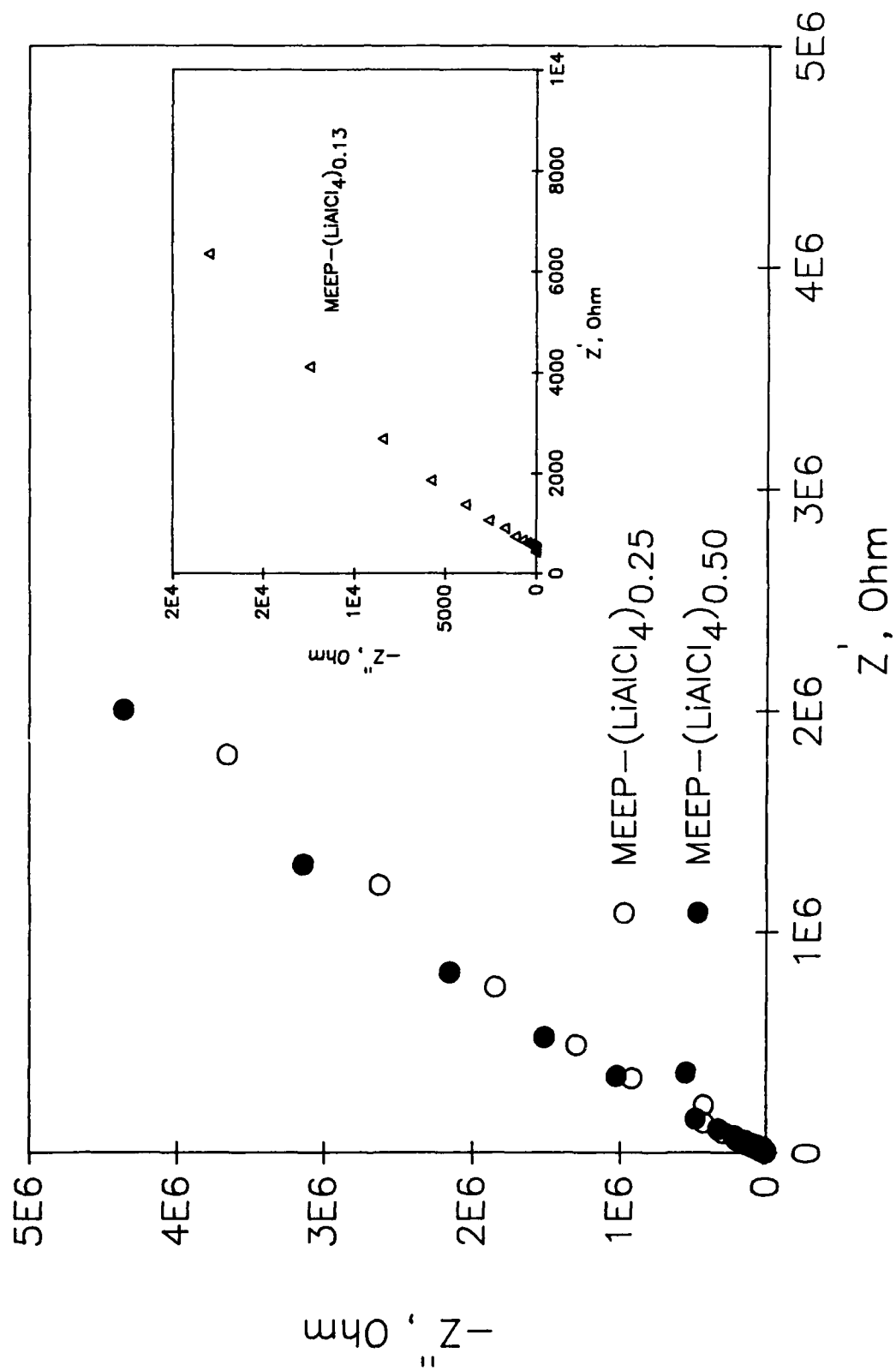


FIGURE 10

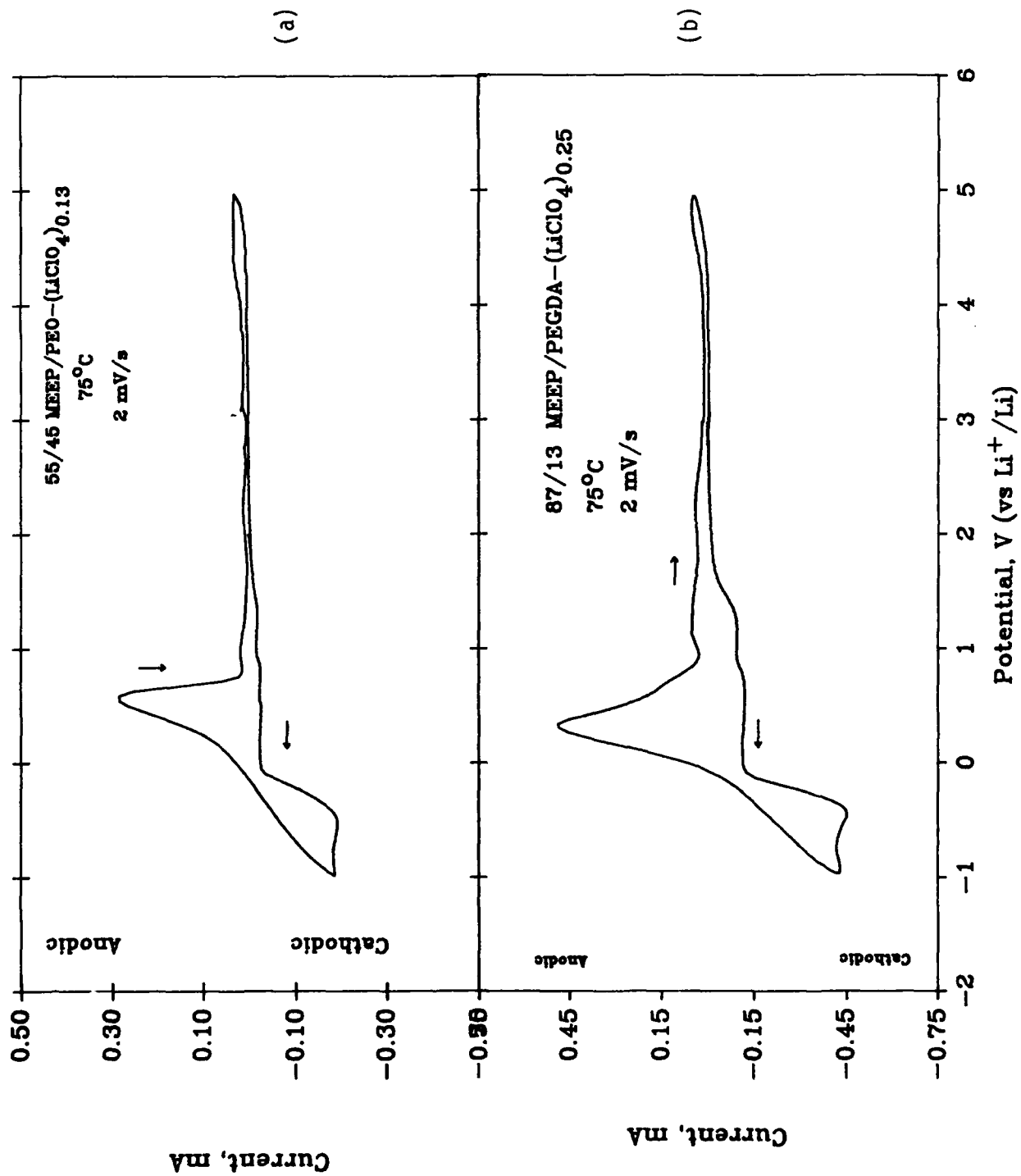


FIGURE 11

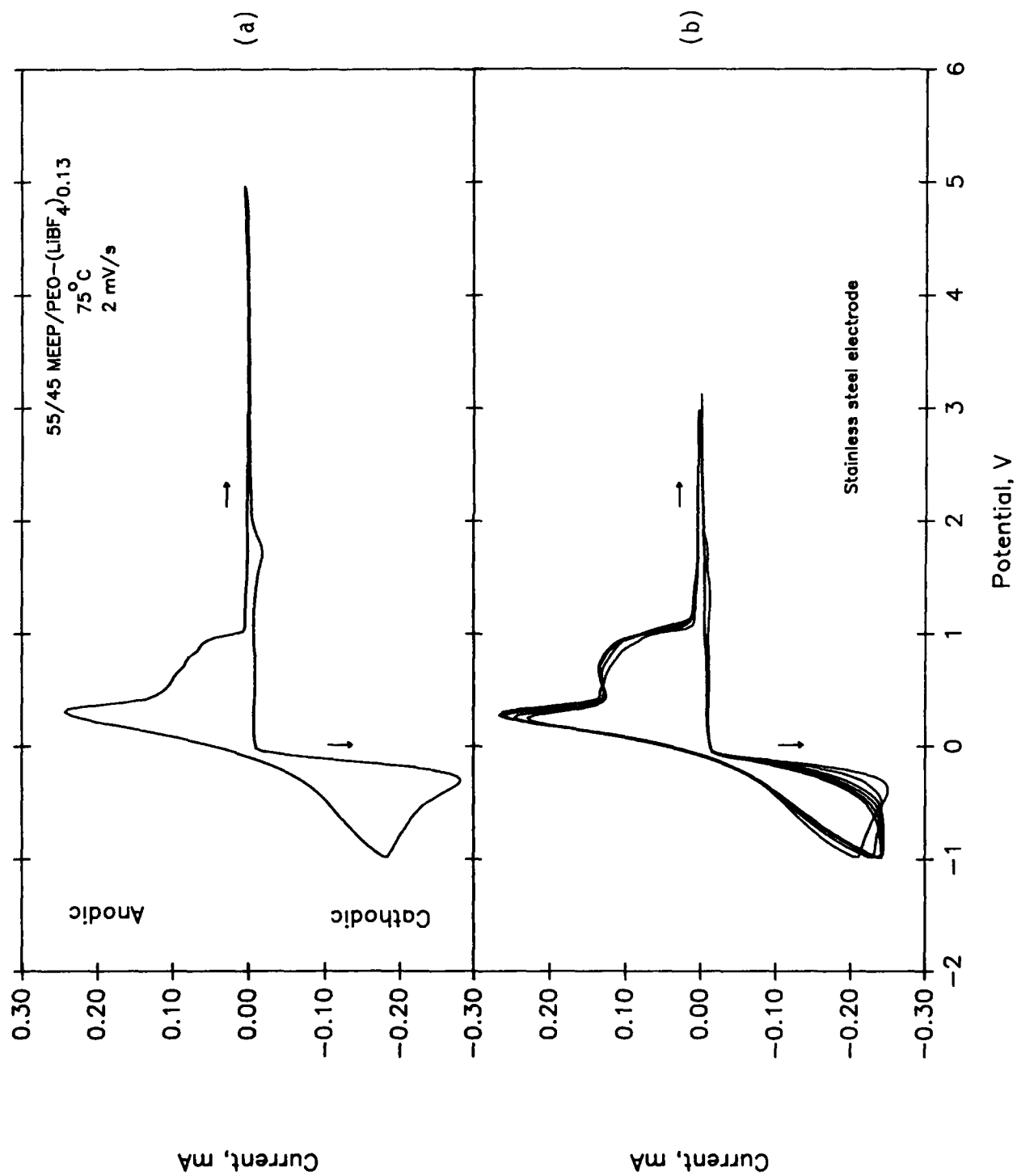


FIGURE 12

Electrolyte: 55 MEEP/45 PEO-(LiClO₄)_{0.13}

Temperature = 75°C

Scan rate = 0.5 mV/s

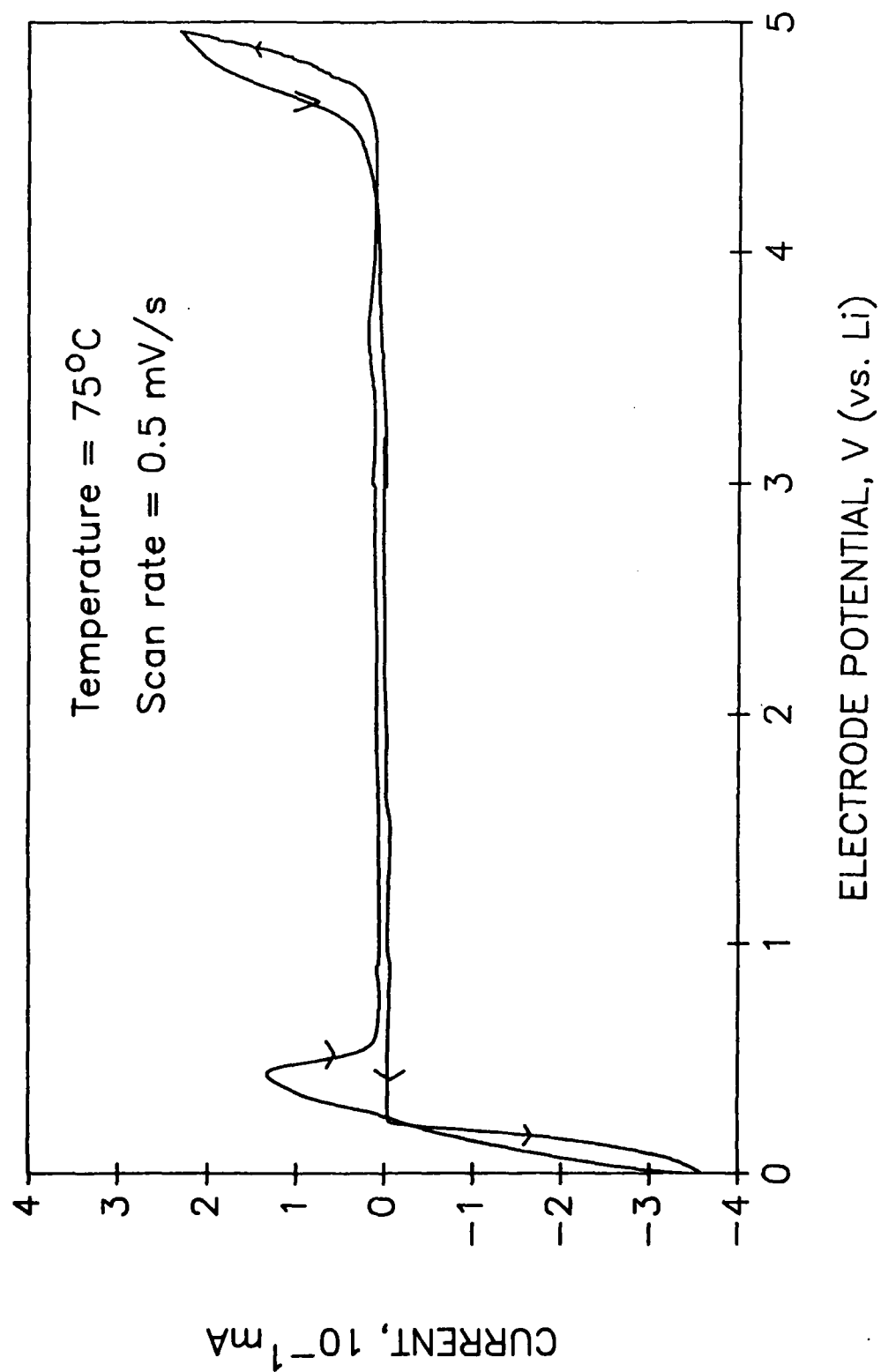


FIGURE 13

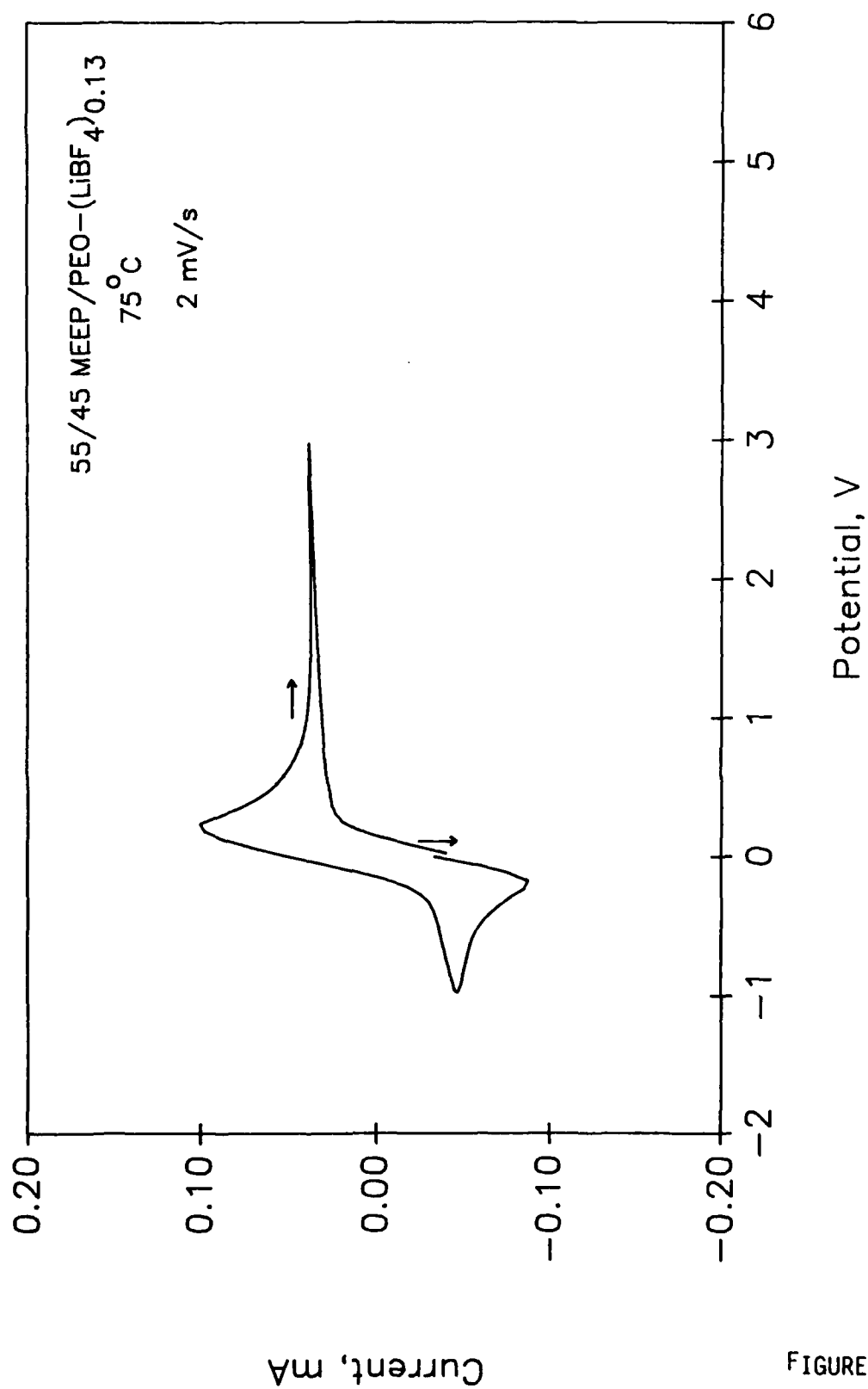


FIGURE 14

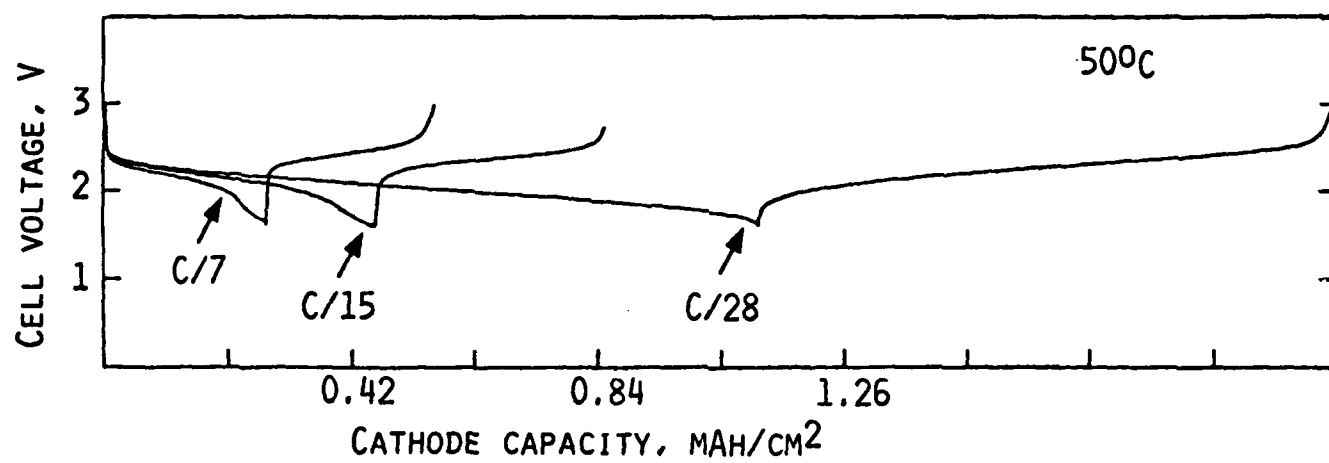


FIGURE 15

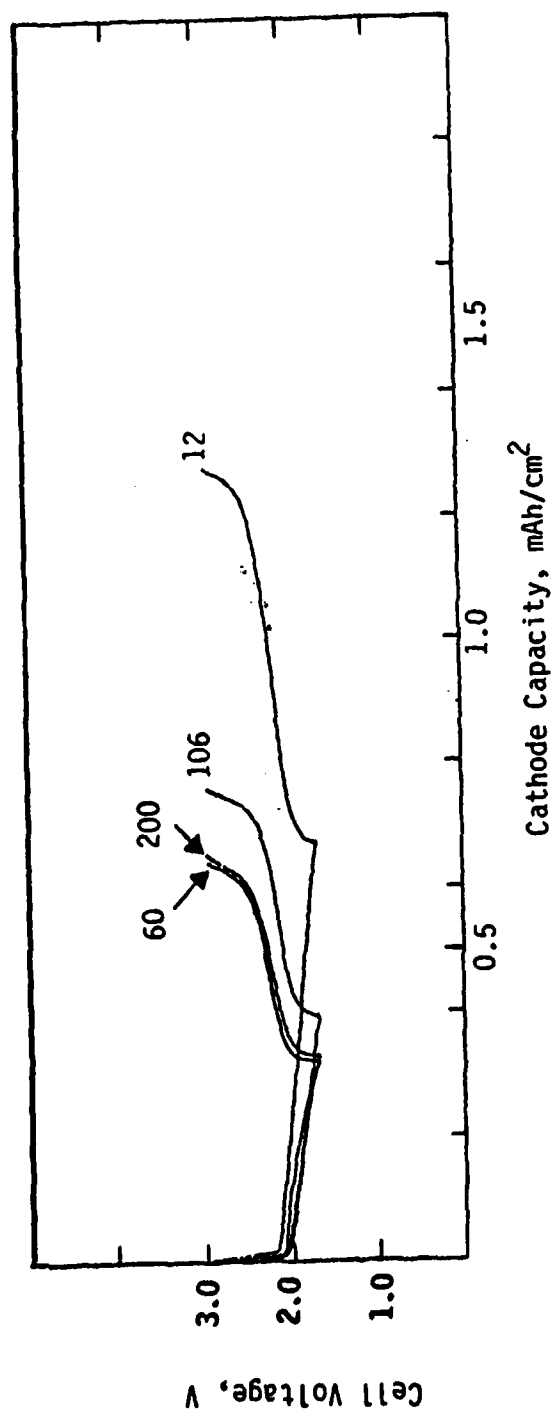


FIGURE 16

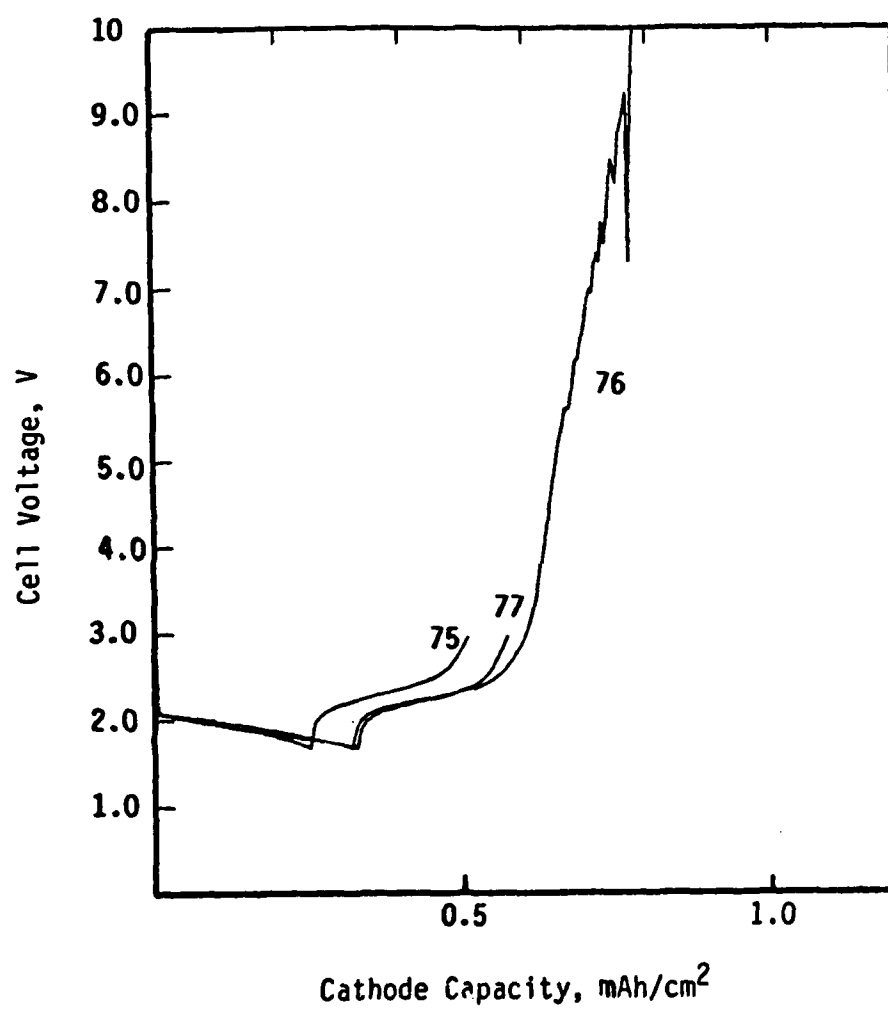


FIGURE 17

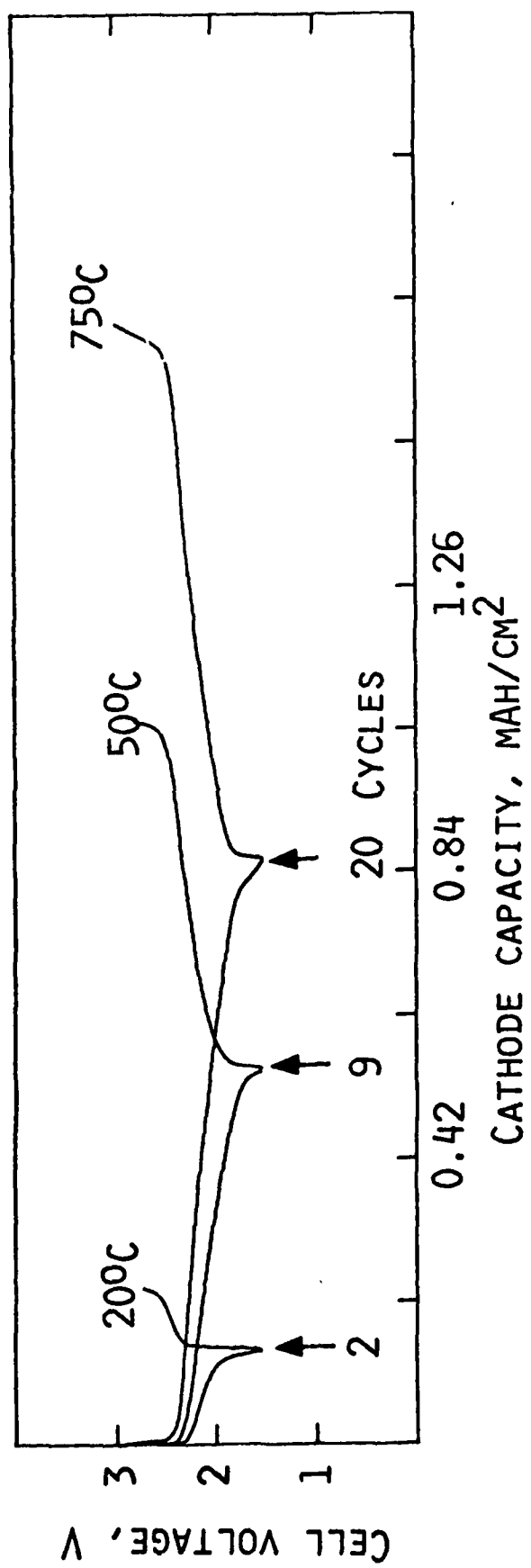


FIGURE 18